

OM of: US-09-303-518D-131 to: A\_Geneseq\_032802.\* out\_format : pfs

Date: Jun 30, 2002 6:41 AM

About: Results were produced by the GenCore software, version 4.5.  
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Command line parameters:

-MODEL=frame+n2p.model -DEV=xlh  
-O=/cgn2.1/USPTO.spool/US09303518/runat\_28062002.142712.4291/app\_query.fasta\_1.23501  
-DB=A\_Geneseq\_032802 -QFMT=Fastan -SUFFIX=rag -GAPOP=12.000  
-GAPEXT=4.000 -MINMATCH=0.100 -LOOPCL=0.000 -LOPEXT=0.000  
-GAPOP=4.500 -QGAPEXT=0.050 -XGAPOP=10.000 -XGAPEXT=0.500  
-GAPOP=6.000 -FGAPEXT=7.000 -START=1 -MATRIX=blosum62  
-DELOP=6.000 -DELEXT=7.000 -SCORE=1 -THR\_SCORE=pcr  
-THRX=human40.cdi -LIST=100 -DOCALIGN=200 -THR\_SCORE=pcr  
-THRM=100 -THR\_MIN=0 -ALIGN=45 -MODE=LOCAL -OUTPMT=pfs  
-NORM=ext -HEAPSIZE=500 -MINLEN=0 -MAXLEN=2000000000  
-USER=US09303518 -CGNL\_1.0 -NCPU=6 -ICPU=3 -LONGLOG  
-DEV\_TIMEOUT=120 -WARN\_TIMEOUT=30 -NO\_XLPHY -WAIT -THREADS=1.

Search information block:

Query: US-09-303-518D-131

Query length: 1344

Database: A\_Geneseq\_032802.\*

Database sequences: 747574

Database length: 111073796

Search time (sec): 627.340000

score list:

Sequence	Strd Orig	ZScore	EScore Len	Documentation
/SIDSL1/gcgdata/geneseq/geneseq-emb1/AA1999.DAT:AAU39645	44	3.9e-237	2281.00	4391.99
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/SIDSL1/gcgdata/geneseq/geneseq-emb1/AA1999.DAT:AAU38563	32	5.5e-167	1631.00	3135.97
/SIDSL1/gcgdata/geneseq/geneseq-emb1/AA1999.DAT:AAU38564	44	1.0e-72	757.00	1448.34
/SIDSL1/gcgdata/geneseq/geneseq-emb1/AA1999.DAT:AAU38565	45	1.1e-60	648.00	1223.88
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/SIDSL1/gcgdata/geneseq/geneseq-emb1/AA1999.DAT:AAU38569	44	2.5e-39	450.50	840.30
/SIDSL1/gcgdata/geneseq/geneseq-emb1/AA1999.DAT:AAU38570	44	2.5e-38	438.00	816.21
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/SIDSL1/gcgdata/geneseq/geneseq-emb1/AA2001.DAT:AAU51862	+	101.50	175.06	0.0742
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/SIDSL1/gcgdata/geneseq/geneseq-emb1/AA2001.DAT:AAU39374	+	101.00	153.50	0.2236
/SIDSL1/gcgdata/geneseq/geneseq-emb1/AA2001.DAT:AAU58082	+	100.50	166.11	0.1328
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seq\_identification\_block:

ID AAY38564 standard; Protein: 447 AA.

XX AAY38564;

XX 08-OCT-1999 (first entry)

DE Neisseria gonorrhoeae antigen encoded by ORF22.

KW Neisseria meningitidis; Neisseria gonorrhoeae; antigen; vaccine;



384 alMetProLeuAspIleLeuProThrLeuLeuLeuLeuArgAspLeuIleVal 400  
 1201 GGCCATACCGACAGCGAGGCTTGGGTTGCTTGGCAATTGGACGAAGA 1250  
 401 GlyAspThrAspSerAlaGlnAlaLeuGlyCysLeuGluLeuAspGlu 417  
 1251 AGACCTCGCTTGTGAGCTTCTGCTGCGCGGCAAAATACGAATAGCGCC 1300  
 417 uAspLeuAlaLeuCysSerPheValCysProGlyLysTyrgLutyrGlyp 434  
 1301 CGCTGTGCGCAAGAGTCTGGAACCATTCAGAGGAAGGC 1341  
 434 roLeuLeuArgLysValLeuGluThrIleGluLysGluGly 447

seq\_name: /SIDSI/gcgdata/geneseq/geneseq-emb1/AA1999.DAT.AAY38561

seq\_documentation\_block:

ID AAY38561 standard; Protein: 447 AA.

AC AAY38561;

DT 08-OCT-1999 (first entry)

Neisseria meningitidis antigen encoded by ORF22.

Neisseria meningitidis; Neisseria gonorrhoeae; antigen; vaccine;  
 treatment; Neisseria infection; meningitis; septicaemia; gonorrhea.

Neisseria meningitidis.

W09924578-A2.

20-MAY-1999.

09-OCT-1998; 98WO-IB01665.

01-SEP-1998; 98GB-0019016.

06-NOV-1997; 97GB-0023516.

14-NOV-1997; 97GB-0024190.

18-NOV-1997; 97GB-0024386.

27-NOV-1997; 97GB-0025158.

10-DEC-1997; 97GB-0026147.

14-JAN-1998; 98GB-0000759.

(CHIR-) CHIRON SPA.

Grandi G, Masignani V, Pizza M, Rappuoli R, Scarlato V;

WPI; 1999-327407/27.

N-PSDB; AA212026.

Proteins from Neisseria meningitidis and N. gonorrhoeae useful for  
 diagnosis, treatment and prevention of infection

Claim 4; Page 123; 524pp; English.

Amino acid sequences AAY38499-Y38944 represent Neisseria meningitidis  
 and N. gonorrhoeae antigenic proteins. They are encoded by open  
 reading frames (ORFs) AA211972-Z12358. The antigenic proteins,  
 their fragments, their nucleic acids and antibodies are used for  
 diagnosis, prevention (as vaccines) or treatment of Neisseria  
 infections, such as meningitis, septicaemia and gonorrhea. Both  
 organisms are closely related. Fragments of the nucleic acids  
 are useful as hybridisation probes and antisense reagents.

Sequence 447 AA;

alignment\_scores:

Quality: 2221.00 Length: 447  
 Ratio: 4.980 Gaps: 0  
 Percent Similarity: 99.776 Percent Identity: 96.197

alignment\_block:

US-09-303-518D-131 x AAY38561 ..

Align seg 1/1 to: AAY38561 from: 1 to: 447

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 1 MetIleLysIleLysLysGlyLeuAsnLeuProIleAlaGlyArgProG 17  
 51 GCAAGTCATTTATGACGGCCGCGCATTTACCGAAAGTCGCGTTGTTGGCG 100  
 17 uGlnAlaValTyrAspGlyProAlaIleThrGluValAlaLeuLeuGly 34  
 101 AAGATATGTCGGCATCGGCCCTCGATGAAATCAAGGAAGTGAAGCC 150  
 34 LuGluTyrAlaGlyMetArgProSerMetLysValLysGluGlyAspAla 50  
 151 GTCAAAAAGGCCAAGTCTGTTTGAAGACAAAAAAGATCCGGCGTAGT 200  
 51 ValLysLysGlyGlnValLeuPheGluAspLysLysAsnProGlyValVa 67  
 201 ATTTACTGCGCGGCTTCAGCAAAATCGCGCTATTACCGTGGCGAAA 250  
 67 lPheThrAlaProAlaSerGlyLysIleAlaAlaIleHisArgGlyGluL 84  
 251 AGCGCGTACTTCAGTCAGTCGTGATTGCCGTGAAGGCAACGACGAATC 300  
 84 yArgValLeuGlnSerValIleAlaValGluGlyAsnAspGluIle 100  
 301 GAGTTCGAAGCTACGTACCTGGAAGCGCTGGCAAAATTGAGCAGCGAAA 350  
 101 GluPheGluArgTyrAlaProGluAlaLeuAlaAsnLeuSerGlyGluG 117  
 351 AGTCGCGCGCAACTGATTCAATCAGGCTTATGACTGCGCTTCGCACCC 400  
 117 uValArgArgAsnLeuIleGlnSerGlyLeuTrpThrAlaLeuArgThra 134  
 401 GTCGTTTCAGCAAAATCCCTGCGGTAGATGCCGAGCGGTTCGCCATCTTC 450  
 134 rgProPheSerLysIleProAlaValAspAlaGluProPheAlaIlePhe 150  
 451 GTCAATGCGATGAGCACCAATCCGCTGCGGACCTACGGTGCATCAT 500  
 151 ValAsnAlaMetAspThrAsnProLeuAlaAlaAspProThrValIle 167  
 501 CAAAGAGCGCGCGAAGACTTCAACGCGGCTGTTGTTGATTGAGCGGCC 550  
 167 eLysGluAlaAlaGluAspPheLysArgGlyLeuLeuValLeuSerArgL 184  
 551 TGACCGAAGCGTAAATCCATGTGTAAAGCAGCAGCGCGACGACGTCGCG 600  
 184 euThrGluArgLysIleHisValCysLysAlaAlaGlyAlaAspValPro 200  
 601 TCTGAAATGCTGCCAATATCGAAACACATGAATTTGGCGGCGCCGATCC 650  
 201 SerGluAsnAlaAlaAsnIleGluThrHisGluPheGlyGlyProHisPr 217  
 651 TGCGCGCTTGAGTGGCAGCACACATTCATTCATTCGAGCAGTCGCGCGGA 700  
 217 oAlaGlyLeuSerGlyThrHisIleHisPheIleGluProValGlyAlaA 234  
 701 ATAAACCGGTGGACCATCAATATCAAGACGTGATTGCTATCGGACGT 750  
 234 snLysThrValTrpThrIleAsnTyrGlnAspValIleThrIleGlyArg 250  
 751 TTGTTGCTAACAGCGCTCTGAATACCGCGCGTGGTTCCTTGGCGGG 800  
 251 LeuPheAlaThrGlyArgLeuAsnThrGluArgValIleAlaLeuGlyG 267  
 801 CCTGCAAGTCAACAAACCGCGCTCTTGGCTACCGTTCCTTGGTGGGAAGG 850  
 267 ySerGlnValAsnLysProArgLeuLeuArgThrValLeuGlyAlaLysV 284





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67 lPheThrAlaProAlaSerGlyLysIleAlaAlaIleHisArgGlyGluL 84
251 AGCGCGTACTTCACTCAGTCTGATTCGCGTTGAAGGCAACGACGAAATC 300
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84 ysArgValLeuGlnSerValValIleAlaValGluGlyAsnAspGluIle 100
301 GAGTTCGACGCTACGTACCTGAGCTGAGCGTGGCAAAATTTGAGCAGCGAAA 350
|||||
101 GluPheGluArgTyrValProGluAlaLeuAlaLysLeuSerSerGluLy 117
351 AGTGCAGCGCAACCTGATTCACCTCAGCTTATGACTGCGCTTCGCAACC 400
|||||
117 sValArgGlnLeuGlnSerGlyLeuThrAlaLeuArgThrA 134
401 GTCCGTTTCAGCAAAATCCCTGCGGTAGATCCGAGCGTTCGCACTTC 450
|||||
134 rgProPheSerLysIleProAlaValAspAlaGluProPheAlaIlePhe 150
451 GTCATCGGATGACACCAATCCGCTGGCTGCCGACCTACGCTCATCAT 500
|||||
151 ValAsnAlaMetAspThrAsnProLeuAlaAlaAspProThrValIleIl 167
501 CAAGAAGCGCGCAAGACACTTCAACCGCGCTGTTGTTGATTTAGCGGCC 550
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167 eLysGluAlaAlaGluAspPheLysArgGlyLeuLeuValLeuSerArgL 184
551 TGACCGAAGCTTAAATCCATGTGTGTAAGCAGCAGCGCAGAGTCCCG 600
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184 euThrGluArgLysIleHisValCysLysAlaAlaGlyAlaAspValPro 200
601 TCTGAAATGCTGCAATATCGAACAACATGATTTGGCGCGCGCATCC 650
|||||
201 SerGluAsnAlaAlaAsnIleGluThrHisGluPheGlyGlyProHisPr 217
651 TGCCTGCTGAGTGGCGACGACATTCATTCATCGACCGCAGTGGCGCGA 700
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217 oAlaGlyLeuSerGlyThrHisIleHisPheIleGluProValGlyAlaA 234
701 ATAAACCGTGTGACCATCAATATCAAGACGTGATTCGTCGAGCGT 750
|||||
234 snLysThrValTrpThrIleAsnTyrGlnAspValIleAlaIleGlyArg 250
751 TTGTTCTGTAACAGCGCGCTGAATACCGAGCGGTGTTGCTTGGCGG 800
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251 LeuPheValThrGlyArgLeuAsnThrGluArgValValAlaLeuGlyG 267
801 COTGCAAGTCAACAACCGCGCTCTTCCGTACCGTGGTGGTGGCGAAG 850
|||||
267 yLeuGlnValAsnLysProArgLeuLeuArgThrValLeuGlyAlaLysV 284
851 TCTCTCACTTACCGCGCGCAATTTGTTGACGCGCAACCGCGTGTATT 900
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284 alSerGlnLeuThrAlaGlyGluLeuValAspAlaAspAsnArgValIle 300
901 TCCGTTCCGTTATGACCGGTGCGATTCACAAAGCGCGCATGATTATT 950
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301 SerGlySerValLeuAsnGlyAlaIleAlaGlnGlyAlaHisAspTyrLe 317
951 GGGAGCTACCAAT 966
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317 uGlyArgTyrHisAsn 322
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seq_documentation_block:
ID AA1999 standard; protein: 158 AA.
XX
AC AA1999;
XX
DT 08-OCT-1999 (first entry)
XX
DE Neisseria meningitidis antigen encoded by a partial ORF22.
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XX
KW Neisseria meningitidis; Neisseria gonorrhoeae; antigen; vaccine;
PN treatment; Neisseria infection; meningitis; septicaemia; gonorrhea.
XX Neisseria meningitidis.
XX WO9924578-A2.
XX
PD 20-MAY-1999.
XX
PF 09-OCT-1998; 98WO-IB01665.
XX
PR 01-SEP-1998; 98GB-0019016.
PR 06-NOV-1997; 97GB-0023516.
PR 14-NOV-1997; 97GB-0024190.
PR 18-NOV-1997; 97GB-0024386.
PR 27-NOV-1997; 97GB-0025158.
PR 10-DEC-1997; 97GB-0026147.
PR 14-JAN-1998; 98GB-0000759.
XX
PA (CHIR-) CHIRON SPA.
XX
PI Grandi G, Masignani V, Pizza M, Rappuoli R, Scarlato V;
XX
DR WPI; 1999-327407/27.
DR N-PSDB; AA12025.
XX
PT Proteins from Neisseria meningitidis and N. gonorrhoeae useful for
PT diagnosis, treatment and prevention of infection
XX
PS Claim 4; Page 122; 524pp; English.
XX
CC Amino acid sequences AAY38499-Y38944 represent Neisseria meningitidis
CC and N. gonorrhoeae antigenic proteins. They are encoded by open
CC reading frames (ORFs) AA11972-212358. The antigenic proteins,
CC their fragments, their nucleic acids and antibodies are used for
CC diagnosis, prevention (as vaccines) or treatment of Neisseria
CC infections, such as meningitis, septicaemia and gonorrhea. Both
CC organisms are closely related. Fragments of the nucleic acids
CC are useful as hybridisation probes and antisense reagents.
XX
SQ Sequence 158 AA;
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alignment_scores:
Quality: 757.00 Length: 158
Ratio: 4.822 Gaps: 0
Percent Similarity: 99.367 Percent Identity: 93.671

alignment_block:
US-09-303-518D-131 x AAY38560 ..

Align seg 1/1 to: AAY38560 from: 1 to: 158

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1 MetIleLysIleLysLysGlyLeuAsnLeuProIleAlaGlyArgProG 17
51 GCAAGTCATTTATGACGCGCGCGCATTCAGGAGTCGCGTCTCTGGCG 100
|||||
17 uGlnAlaValTyrAspGlyProAlaIleThrGluValAlaLeuLeuGlyG 34
101 AAGAAATATGTCGCGATCGCGCTCGATGAAATCAAGGAAGGTGAAGC 150
|||||
34 luGluTyrAlaGlyMetArgProSerMetLysValLysGluGlyAspAla 50
151 GTCAAAAAGCGCAAGTCTGTTTGAAGCAAAAAGATCCGGCGCTAGT 200
|||||
51 ValLysLysGlyGlnValLeuPheGluAspLysLysAsnProGlyValVa 67
201 ATTACTGCGCGCGCTTCAGGCAAAATCGCGCTATTTCACCGTGGCGAA 250
|||||
67 lPheThrAlaProAlaSerGlyLysIleAlaAlaIleHisArgGlyGluL 84
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251 AGCGCTACTTCACTCAGTCGTGATTCCGCTTGAAGCGCAACGACGAAATC 300  
 |||||  
 84 ysargValLeuGlnSerValValIleAlaValGlu\*\*AsnAspGluIle 100  
 |||||  
 301 GAGTTCGACGCTACTGACCTGAGCGCTGGCAAAATTCAGCAGCGAAA 350  
 |||||  
 101 GluPheGluArgTyrAlaProGluAlaLeuAlaAsnLeuSerGlyGluG 117  
 |||||  
 351 AGTGGCCCGCAACCTGATCAATCAGGCTTATGGACTGCGCTTCGACCC 400  
 |||||  
 117 uValArgArgAsnLeuIleGlnSerGlyLeuTrpThrAlaLeuArgThra 134  
 |||||  
 401 GTCCGCTCAGCAAAATCCCTGCGTAGATGCCGAGCGCTTCGCCATCTTC 450  
 |||||  
 134 rgProPheSerLysIleProAlaValAspAlaGluProPheAlaIlePhe 150  
 |||||  
 451 GTCATCGCATGGACACCAATCCG 474  
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 151 ValAsnAlaMetAspThrAsnPro 158

seq\_name: /SIBS1/gcgdata/geneseq/geneseq-emb1/AA1999.DAT:AAV34439

seq\_documentation\_block:

ID AAY34439 standard; Protein; 451 AA.

AC AAY34439;

XX  
 XX  
 XX 25-AUG-1999 (first entry)

XX Porphyromonas gingivalis protein PGI.

XX Porphyromonas gingivalis; PG; periodontal disease; gingivitis;  
 KW vaccine; antigenic.

XX Porphyromonas gingivalis.

XX OS

XX PN WO9229870-A1.

XX PD 17-JUN-1999.

XX PF 10-DEC-1998; 98WO-AU01023.

XX PR 04-AUG-1998; 98AU-0005028.

XX PR 10-DEC-1997; 97AU-0000839.

XX PR 31-DEC-1987; 97AU-0001182.

XX PR 30-JAN-1998; 98AU-0001546.

XX PR 10-MAR-1998; 98AU-0002264.

XX PR 09-APR-1998; 98AU-0002911.

XX PR 23-APR-1998; 98AU-0003128.

XX PR 05-MAY-1998; 98AU-0003338.

XX PR 22-MAY-1998; 98AU-0003654.

XX PR 29-JUL-1998; 98AU-0004917.

XX PA (CSLC-) CSL LTD.

XX PI Agius CT, Barr IG, Hocking DM, Margetts MB, Patterson MA;

XX PI Ross BC, Rothel LJ, Webb EA;

XX DR WPI; 1999-385613/32.

XX DR N-PSDB; AAX91657.

XX PT Antigenic Porphyromonas gingivalis peptides for preventing

XX PT gingivitis

XX PS Claim 1; Page 417-418; 588pp; English.

XX CC AAX91536 to AAX91801 encode two hundred and sixty six antigenic  
 CC Porphyromonas gingivalis (PG) polypeptide sequences given in AAY34318 to  
 CC AAY34583. AAX91802 to AAX91989 represent PCR primers used in the  
 CC isolation of the PG polypeptides. The PG polypeptides have antibacterial  
 CC activity with a vaccine mechanism of action. The PG polypeptides can be  
 CC used as vaccines especially against Porphyromonas gingivalis. Probes can

CC be used to detect Porphyromonas gingivalis in standard hybridisation  
 CC assays. Porphyromonas gingivalis is involved in periodontal disease  
 CC especially gingivitis.

XX  
 SQ Sequence 451 AA;

alignment\_scores:

Quality: 648.00 Length: 452

Ratio: 2.189 Gaps: 7

Percent Similarity: 65.487 Percent Identity: 34.071

alignment\_block:

US-09-303-518D-131 x AAY34439 ..

Align seg 1/1 to: AAY34439 from: 1 to: 451

1 ATGATTAATAAATCAAAAAAGTCTAAATCTGCCCATCGCGGACACCGGA 50  
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 4 ValIleLysThrLysLysGlyLeuAlaLeuAsnLeuLysGlyLysProLe 20  
 :|||  
 51 GCAAGTCATTTATGACGCGCCGCTTACCGAAGTC...GCCTTGCTTG 97  
 :|||  
 20 uProGluMetLeuAlaGluProAlaGlnSerProThrTyrAlaValValp 37  
 :|||  
 98 GCGAAGAATATGTCGCGCATGCGCCCTCGATGAAATCAAGGAGGTGAA 147  
 :|||  
 37 roAspAspPheGluGlyValIleProLysValThrAlaArgProGlyAsp 53  
 :|||  
 148 GCCGTCAAAAAAGCCAAAGTCTGTTTGAAGACAAAAAAGAAATCCGGCGT 197  
 :|||  
 54 LysValArgAlaGlySerAlaLeuMetHisHisLysAlaTyrProGluMe 70  
 :|||  
 198 AGTATTTACTGCGCGCTTCAGGCAAAATCGCGCTTATTCACCGTGGCG 247  
 :|||  
 70 tLysPheThrSerProValSerGlyGluValIleAlaValAsnArgGlyA 87  
 :|||  
 248 AAAAGCGCTACTTCAGTCAGTCGTGATTCGCTTGAAGGCAACGACGAA 297  
 :|||  
 87 laLysArgLysValLeuSerIleGluValLysProAspGlyLeuAsnGlu 103  
 :|||  
 298 ATCGAG...TTCGAACGCTACGTACCTGAGCGCTGGCAAAATTCAGCAG 344  
 :|||  
 104 TyrGluSerPheProValGlyAspProSerAla.....LeuSerAl 117  
 :|||  
 345 CGAAAAAGTCGCGCGCAACCTGATTCATCAGCTTATGGACTGCGCTTC 394  
 :|||  
 117 aGluGlnIleLysGluLeuLeuSerSerGlyMetTrpGlyPheIleLe 134  
 :|||  
 395 GCACCCGCTCGCTTCAGCAAAATCCCTGCGGTAGATCGCGCGCTTCGCC 444  
 :|||  
 134 ysGlnArgProTyrAspIleValAlaThrProAspIleAlaProArgAsp 150  
 :|||  
 445 ATCTTCGTCATCGGATGACACCAATCCGCTGCGCGACCCCTACGCT 494  
 :|||  
 151 IleTyrIleThrAlaAsnPheThrAlaProLeuAlaProAspPheAspPh 167  
 :|||  
 495 CATCATCAAGAAGCGCGCGAGACTTCAACGCGCTGTTGGTATTCA 544  
 :|||  
 167 eIleValArgGlyGluGluArgAlaLeuGlnThrAlaIleAspAlaLeu 184  
 :|||  
 545 GCGCGCTGACCGCAACGTAATCCATGTGTGTAAGCAGCAGCGCGCAGAC 594  
 :|||  
 184 laLysLeuThrThrGlyLysValTyrValGlyLeuLysProGlySerSer 200  
 :|||  
 595 GTGCGCTCTGAAATGCTCCCAATATCCAAACACATGAATTTGGCGGCC 644  
 :|||  
 201 LeuGlyLeuHisAsnAlaGluIleValGluValHis.....GlyPr 214  
 :|||  
 645 GCATCTCTGCGCGCTTGTAGTGGCGCAGCATTCATTCATCGAGCAGTCG 694  
 :|||  
 214 ohisProAlaGlyAsnValGlyValIleAsnHisThrLysProIleA 231



```

695 GCGGATAAACCCTGGACCATCAATTATCAAGACGTGATTGCTATC 744
:: : : : : : : : : : : : : : : : : : : : : : : : : :
231 snArgGlyGluThrValTrpThrLeuLysAlaThrAspLeuIleValIle 247
:: : : : : : : : : : : : : : : : : : : : : : : : : :
745 GGAGCTTTGTTCTGAACACCGCTCTGAATACCGAGCGGTGGTTCCTT 794
:: : : : : : : : : : : : : : : : : : : : : : : : : :
248 GlyArgPheLeuLeuThrGlyLysAlaAspPheThrArgMetIleAlaMe 264
:: : : : : : : : : : : : : : : : : : : : : : : : : :
795 GGGCGGCTGCAAGTCAACAAACCGCCCTCTTCGCTACCGTTCGGTG 844
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264 tThrGlySerAspAlaAlaHisGlyTyrValArgIleMetProGlyC 281
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845 CGAAGGTCTCACTTACCGCGCGCAATTGGTT.....GACGCGGAC 888
:: : : : : : : : : : : : : : : : : : : : : : : : : :
281 yAsnValPheAlaSerPheProGlyArgLeuThrIleLysGluSerHis 297
:: : : : : : : : : : : : : : : : : : : : : : : : : :
889 AACCGCTGATTTCGGTTCGGTATTGAACGGTTCGATTGCACAAGCGC 938
:: : : : : : : : : : : : : : : : : : : : : : : : : :
298 GluArgValIleAspGlyAsnValLeuThrGlyLysLysLeuCysGluLy 314
:: : : : : : : : : : : : : : : : : : : : : : : : : :
939 GCATGATTATTGGACGCTACCAACATCAGATTTCGGTTCGAGAAG 988
:: : : : : : : : : : : : : : : : : : : : : : : : : :
314 sGluProPheLeuSerAlaArgCysAspGlnIleThrValIleProGlu 331
:: : : : : : : : : : : : : : : : : : : : : : : : : :
989 GCCGAGC...AAAGAGCTGTTCGGCTGGTTCGCGCGACCGGACAAA 1035
:: : : : : : : : : : : : : : : : : : : : : : : : : :
331 lyAspValAspGluLeuPheGlyTrpAlaAlaProArgLeuAspGln 347
:: : : : : : : : : : : : : : : : : : : : : : : : : :
1036 TACTCCATCAGCGCACCTCTCGGCCATTTCCTA...AAACAAACT 1082
:: : : : : : : : : : : : : : : : : : : : : : : : : :
348 TyrSerMetSerArgAlaTyrPheSerTrpLeuGlnGlyLysAsnLysG 364
:: : : : : : : : : : : : : : : : : : : : : : : : : :
1083 CTCAAGTTCAGCAGCGGCAACGCGGCGGCGGCGGCGGCGGCGGCGG 1132
:: : : : : : : : : : : : : : : : : : : : : : : : : :
364 uTyrValLeuAspAlaArgIleLysGlyGlyGluArgAlaMetIleMetS 381
:: : : : : : : : : : : : : : : : : : : : : : : : : :
1133 TCGCAGCTATTAGCGCGGTAAATCGGTGGACATCTCGCTACCTTGC 1182
:: : : : : : : : : : : : : : : : : : : : : : : : : :
381 eAsnGluTyrAspArgValPheProMetAspIleTyrProGluTyrLeu 397
:: : : : : : : : : : : : : : : : : : : : : : : : : :
1183 TTGCGCGATTAAATCGTCGGCGATACGACGCGGCGGCGGCGGCGGTT 1232
:: : : : : : : : : : : : : : : : : : : : : : : : : :
398 LeuLysAlaIleIleAlaPheAspLleAspLysMetGluAspLeuGlyI 414
:: : : : : : : : : : : : : : : : : : : : : : : : : :
1233 CTGGAATTGGACGAAGACCTCGCTTTGTCAGCTTCGCTGCGCGG 1282
:: : : : : : : : : : : : : : : : : : : : : : : : : :
414 eTyrGluValAlaProGluAspPheAlaThrCysGluPheValAspThrS 431
:: : : : : : : : : : : : : : : : : : : : : : : : : :
1283 GCATATAGATACGCGCGGCTGTTCGCAAGTCTGCTGGAACCATTCAG 1332
:: : : : : : : : : : : : : : : : : : : : : : : : : :
431 eRlySileGluLeuGlnArgIleValArgGluGlyLeuAspMetLeuTyr 447
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seq\_name: /SIDSL/gcdata/geneseq/geneseq-emb1/AA1999.DAT:AAAY34318

seq\_documentation\_block:

ID AAY34318 standard; Protein; 454 AA.

XX

AC AAY34318;

XX

25-AUG-1999 (first entry)

XX

Porphyromonas gingivalis protein PGI.

XX

Porphyromonas gingivalis; PG; periodontal disease; gingivitis;

XX

vaccine; antigenic.

XX

Porphyromonas gingivalis.

XX

W09929870-A1.

XX

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XX 17-JUN-1999.
PD
XX 10-DEC-1998; 98WO-AU01023.
PF
XX 04-AUG-1998; 98AU-0005028.
PR 10-DEC-1997; 97AU-0000839.
PR 31-DEC-1997; 97AU-0001182.
PR 30-JAN-1998; 98AU-0001546.
PR 10-MAR-1998; 98AU-0002364.
PR 09-APR-1998; 98AU-0002911.
PR 23-APR-1998; 98AU-0003128.
PR 05-MAY-1998; 98AU-0003338.
PR 22-MAY-1998; 98AU-0003654.
PR 29-JUL-1998; 98AU-0004917.
XX (CSLC-) CSL LTD.
PA
XX Agius CT, Barr IG, Hocking DM, Margetts MB, Patterson MA;
PI Ross BC, Rothel LJ, Webb EA;
XX
XX WPI: 1999-385613/32.
DR N-PSDB; AAX91536.
XX
XX Antigenic Porphyromonas gingivalis peptides for preventing
PT gingivitis
XX
XX Claim 1; Page 277; 588pp; English.
XX
XX AAX91536 to AAX91801 encode two hundred and sixty six antigenic
CC Porphyromonas gingivalis (PG) polypeptide sequences given in AAY34318 to
CC AAY34583. AAX91802 to AAX91989 represent PCR primers used in the
CC isolation of the PG polypeptides. The PG polypeptides have antibacterial
CC activity with a vaccine mechanism of action. The PG polypeptides can be
CC used as vaccines especially against Porphyromonas gingivalis. Probes can
CC be used to detect Porphyromonas gingivalis in standard hybridisation
CC assays. Porphyromonas gingivalis is involved in periodontal disease
CC especially gingivitis.
XX
XX Sequence 454 AA;

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alignment\_scores:

Quality: 648.00 Length: 452

Ratio: 2.189 Gaps: 7

Percent Similarity: 65.487 Percent Identity: 34.071

alignment\_block:

US-09-303-518D-131 x AAY34318 ..

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:::||||| ||||| ||| ::| |||::|

7 ValIleLysThrLysLysGlyLeuAlaLeuAsnLeuLysGlyLysProLe 23

:::||||| ||||| ||| ::| |||::|

51 GCAAGTCATTATGACGGCGCCGCCATTACCGAAGTC...GCGTTCGTTG 97

::: ||||| ::| ::| ::| ::| ::| ::| ::| ::| ::| ::|

23 uProGluMetLeuAlaGluProAlaGlnSerProThrTyrAlaValAlp 40

::: ||||| ::| ::| ::| ::| ::| ::| ::| ::| ::| ::|

98 GCGAAGAAATATGTCGCATCGCCCTCGATGAAATCAAGGAAGTGAA 147

::: ||||| ::| ::| ::| ::| ::| ::| ::| ::| ::| ::|

40 roAspAspPheGluGlyValIleProLysValThrAlaArgProGlyAsp 56

::: ||||| ::| ::| ::| ::| ::| ::| ::| ::| ::| ::|

148 GCCGTCAAAAGGCCAAGTCGTGTTGAAGACAAAAGAAATCCGGCGCT 197

::: ||||| ::| ::| ::| ::| ::| ::| ::| ::| ::| ::|

57 LysValArgAlaGlySerAlaLeuMetHisHisLysAlaTyrProGluMe 73

::: ||||| ::| ::| ::| ::| ::| ::| ::| ::| ::| ::|

198 AGTATTACTGCGCGCTTCAGGCAAAATCGCGCTATTACCGTGCGG 247

::: ||||| ::| ::| ::| ::| ::| ::| ::| ::| ::| ::|

73 tLysPheThrSerProValSerGlyGluValIleAlaValAsnArgGlyA 90

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248 AAAAGCGCGTACTTCAGTCAGTCGCTGATTGCCGTTGAAGCAACACGAA 297



90 laYsArgLYsValLeuSerIleGluValLysProAspGlyLeuAsnGlu 106  
 298 ATCGAG...TTTCAACGCTACGTACCTGAAGCGCTGCGAAAATTTAGCAG 344  
 107 TyrGluSerPheProValGlyAspProSerAla.....LeuSerAl 120  
 345 CGAAAAAGTCGGCCGCAACCTGATTCAATCAGCCTTATGGACTGGCGTTC 394  
 120 aGluInIleLysGluLeuLeuSerSerGlyMetTrpGlyPheIleL 137  
 395 GCACCGCTCGGTTTCAGCAAAATCCCTGCCGTAGATCCGAGCGGTTCGC 444  
 137 ysGlnArgProTyrAspIleValAlaThrProAspIleAlaProArgasp 153  
 445 ATCTTCGTCAATGCGATGGACACATTCGCTGGCTGCGACCCCTACCGT 494  
 154 IleTyrIleThrAlaAsnPheThrAlaProLeuAlaProAspPheAspPh 170  
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 170 eileValargGlyGluGluargAlaLeuGlnThrAlaIleAspAlaLeuA 187  
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 595 GTGCGGTCTCAAAATGCTCCAATATCGAAACACATGAATTTGGCGGCC 644  
 204 LeuGlyLeuHisAsnAlaGluIleValGluValHis.....GlyPr 217  
 645 GCATCTCGCGGCTTGAGTGGCAGCAGCACATTCATTCATCGAGCCAGTCG 694  
 217 oHisProAlaGlyAsnValcIlyValLeuIleAsnHisThrLysProIleA 234  
 695 GCGCGATAAAACCGTCTGGACCAATCAATATCAAGAGCTGATGCTATC 744  
 234 snArgGlyGluThrValTrpThrLeuLysAlaThrAspLeuIleValIle 250  
 745 GGACGTTGTGTCACAGCGCGCTCTGAATACCGCGCGTGGTGGCTT 794  
 251 GlyArgPheLeuLeuThrGlyLysAlaAspPheThrArgMetIleAlaMe 267  
 795 GGCGGCGCTGCAAGTCACAAACCGCGCTCTGCGTACCGTCTTTGGGTG 844  
 267 tThrGlySerAspAlaAlaHisGlyTyrValArgIleMetProGlyC 284  
 845 CGAAGGTGTCAACTTACCGCGCGCAATTCGGT.....GAGCGCGAC 888  
 284 ysAsnValPheAlaSerPheProGlyArgLeuThrIleLysGluSerHis 300  
 889 AACCGCGTGATTTCGGTTCGGTATTGAACGTCGCGATTGCACAAAGCGC 938  
 301 GluArgValIleAspGlyAsnValLeuThrGlyLysLysLeuCysGluLy 317  
 939 GCATGATTATTTGGACGCGTACCACAAATCAGATTTCGTTATTCGAGAAG 988  
 317 sGluProPheLeuSerAlaArgCysAspGlnIleThrValIleProGlu 334  
 989 GCGCGAGC...AAAGACTGTTTCGGCTGGTGGCGCGCAGCCGACAAA 1035  
 334 LyAspAspValAspGluLeuPheGlyTrpAlaAlaProArgLeuAspGln 350  
 1036 TACTCCATCAGCGACCACTCTCGGCATTTCCTA...AAAACAAACT 1082  
 351 TyrSerMetSerArgAlaTyrPheSerTrpLeuGlnGlyLysAsnLysGl 367  
 1083 CTTCAAGTTCACGACAGCGCTCAACGCGCGCGCAGCCGCTATTCCTA 1132  
 367 uTyrValLeuAspAlaArgIleLysGlyGlyGluArgAlaMetIleMet 384  
 1133 TCGGCACCTATGAGCGCGTATGCGTTGGACATCTCGCTACCTGCCTT 1182

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384 erAsnGluTy rAspArgValPheProMetAspIleTy rProGluTyLeu 400
1183 TTGC GCGATTTAATCTGCGCGATACCGACAGCGCAGGCTTGGGTG 1232
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401 LeuLysAlaIleIleAlaPheAspIleAspIleAspIleMetGluasplLeuGlyII 417
1233 CTTTGGAAATTGGAGGAAGAAGACTCGCTTTGTGTCAGCTTCGTCGCCCG 1282
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417 eTy rGluValAlaIleProGluaspPheAlaThrCysGluPheValAspThrs 434
1283 GC AAATACGAATACGCGCCGCTGTTGCGCAAGTGCTGGAAACCATTGAG 1332
:||| ||| ::::::|||::: |||::: |||:::
434 e rLysIleGluLeuGlnArgIleValArgGluGlyLeuaspMetLeuty r 450
1333 AAGAAA 1338
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451 Lysglu 452
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seq_documentation_block:
ID AAY75271 standard; Protein; 119 AA.
XX AC AAY75271;
XX DT
XX DE 21-MAR-2000 (first entry)
XX KW Neisseria gonorrhoeae ORF 628 protein sequence SEQ ID NO:2016.
XX KW Neisseria meningitidis; Neisseria gonorrhoeae; antigen; vaccine;
XX KW antiseptic; diagnosis; immunogenic; infection; meningitis; septicaemia;
XX OS antibacterial; gene therapy.
XX OS Neisseria gonorrhoeae.
XX PN W09957280-A2.
XX PD
XX PF 11-NOV-1999.
XX PF 30-APR-1999; 99WO-US09346.
XX PR 01-MAY-1998; 98US-0083758.
XX PR 31-JUL-1998; 98US-0094869.
XX PR 02-SEP-1998; 98US-0098994.
XX PR 02-SEP-1998; 98US-0099062.
XX PR 09-OCT-1998; 98US-0103749.
XX PR 09-OCT-1998; 98US-0103794.
XX PR 09-OCT-1998; 98US-0103796.
XX PR 25-FEB-1999; 99US-0121528.
XX PA (CHIR ) CHIRON CORP.
XX PA (GENO-) INST GENOMIC RES.
XX PI Fraser C, Galeotti C, Grandi G, Hickey E, Masignani V, Mora M;
XX PI Petersen J, Pizza M, Rappuoli R, Ratti G, Scalato E, Scarselli M;
XX PI Tettelin H, Venter JC;
XX DR WPI; 2000-062150/05.
XX DR N-PSDB; AA Z54033.
XX XX
XX PT Novel Neisserial polypeptides predicted to be useful antigens for
XX PS vaccines and diagnostics -
XX PS Claim 2; Page 1003; 1453pp; English.
XX CC
XX CC AA Z53015 to AA Z54536, AA Z54577 to AA Z54615, and AA Y74253 to AA Y75941
XX CC represent novel Neisseria meningitis and N. gonorrhoeae polynucleotides
XX CC and polypeptides. AA Z54537 to AA Z54576 and AA Z54616 to AA Z5473 represe
XX CC PCR primers used in the exemplification of the present invention. The
XX CC polypeptides, the polynucleotides, antibodies and compositions of
XX CC the invention can be used as vaccines, as diagnostic reagents, and as
XX CC immunogenic compositions. The polypeptides can be used in the
XX CC manufacture of medicaments for treating or preventing infection due to

```

CC Neisserial bacteria (e.g. meningitis and septicaemia), to detect the  
CC presence of Neisseria bacteria, or to raise antibodies. They may also  
CC be used to screen for agonists or antagonists, which may themselves  
CC have use as antibacterial agents. The polynucleotides of the invention  
CC may also be used in gene therapy protocols.

XX Sequence 119 AA;

alignment\_scores:  
Quality: 582.50 Length: 120  
Ratio: 4.895 Gaps: 1  
Percent Similarity: 99.167 Percent Identity: 99.167

alignment\_block:

US-09-303-518D-131/rev x AAY75271

Align seg 1/1 to: AAY75271 from: 1 to: 119

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1 MetCysValProLeuLysProAlaGlyCysGlyProProAsnSerCysVa 17
|||||
624 TTCATATTCGACGATTTTCAGACGCGCAGCTCTGCGCGCTGCTTTAC 575
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17 lSerIleLeuAlaAlaPheSerAspGlyThrSerAlaProAlaAlaLeuH 34
|||||
574 ACATGATGATTTACGTTGCGTGCAGCGGCTCAATACACAGCGCGGT 525
|||||
34 lThrTrpIleLeuArgSerValArgArgLeuAsnThrAsnArgProArg 50
|||||
524 TTGAAGTCTCGCGGCTCTTTCATGATGACCGTAGGTCGGCAGCGAG 475
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51 LeuLysSerSerAlaAlaSerLeuMetMetThrValGlySerAlaAlaSe 67
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474 CGGATTTGTCCTTCGTCATTCAGCAAGATGCGAAGCGGCTCGCATCTA 425
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67 rGlyLeuValSerIleAlaLeuThrLysMetAlaAsnGlySerAlaSerT 84
|||||
424 CGGAGGATTTTCGTAACGCGGCGGTGCGAAGCGCAGTCCATAAGCCT 375
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84 hrAlaGlyIleLeuLeuAsnGlyArgValArgSerAlaValHisLysPro 100
|||||
374 GATTGAATCAGGTTGCGCGCACATTTTCGCTGCTCAATTTTGCACGCG 325
|||||
101 lAsp...lIleArgLeuArgArgThrPheSerLeuLeuAsnPheAlaSerAl 116
|||||
324 TTCAGGTACG 315
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116 aSerGlyThr 119
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seq\_name: /SIDS1/gcgdata/geneseq/geneseq-emb1/AA2000.DAT:AAAY75272

seq\_documentation\_block:

ID AAY75272 standard; Protein; 120 AA.

XX

AC AAY75272;

XX

DT 21-MAR-2000 (first entry)

XX

DE Neisseria meningitidis ORF 628 protein sequence SEQ ID NO:2018.

XX Neisseria meningitidis; Neisseria gonorrhoeae; antigen; vaccine;  
KW antigenic; diagnosis; immunogenic; infection; meningitis; septicaemia;  
KW antibacterial; gene therapy.

XX

OS Neisseria meningitidis.

XX

PN WO9957280-A2.

XX

PD 11-NOV-1999.

XX

PF 30-APR-1999; 99WO-US09346.

XX 01-MAY-1998; 98US-0083758.  
PR 31-JUL-1998; 98US-0094869.  
PR 02-SEP-1998; 98US-0098994.  
PR 02-SEP-1998; 98US-0099062.  
PR 09-OCT-1998; 98US-0103749.  
PR 09-OCT-1998; 98US-0103794.  
PR 09-OCT-1998; 98US-0103796.  
PR 25-FEB-1999; 99US-0121528.

XX (CHIR ) CHIRON CORP.  
PA (GENO-) INST GENOMIC RES.

XX Fraser C, Galeotti C, Grandi G, Hickey E, Masignani V, Mora M;  
PI Petersen J, Pizza M, Rappuoli R, Ratti G, Scalato E, Scarselli M;  
FI Tettelin H, Venter JC;

XX WPI; 2000-062150/05.

DR N-PSDB; AAZ54034.

XX Novel Neisserial polypeptides predicted to be useful antigens for  
PT vaccines and diagnostics

XX Claim 2; Page 1003; 1453pp; English.

XX AA253015 to AA254536, AA254577 to AA254615, and AAY74253 to AAY75941  
CC represent novel Neisseria meningitis and N. gonorrhoeae polynucleotides  
CC and polypeptides. AA254537 to AA254576 and AA254616 to AA255473 represent  
CC PCR primers used in the exemplification of the present invention. The  
CC polypeptides, the polynucleotides, antibodies and compositions of  
CC the invention can be used as vaccines, as diagnostic reagents, and as  
CC immunogenic compositions. The polypeptides can be used in the  
CC manufacture of medicaments for treating or preventing infection due to  
CC Neisserial bacteria (e.g. meningitis and septicaemia), to detect the  
CC presence of Neisseria bacteria, or to raise antibodies. They may also  
CC be used to screen for agonists or antagonists, which may themselves  
CC have use as antibacterial agents. The polynucleotides of the invention  
CC may also be used in gene therapy protocols.

XX Sequence 120 AA;

alignment\_scores:

Quality: 545.00 Length: 119  
Ratio: 4.698 Gaps: 0  
Percent Similarity: 97.479 Percent Identity: 93.277

alignment\_block:

US-09-303-518D-131/rev x AAY75272

Align seg 1/1 to: AAY75272 from: 1 to: 120

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1 MetCysValProLeuLysProAlaGlyCysGlyProProAsnSerCysVa 17
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624 TTCATATTCGACGATTTTCAGACGCGCAGCTCTGCGCGCTGCTTTAC 575
|||||
17 lSerMetLeuAlaAlaPheSerAspGlyThrSerAlaProAlaAlaLeuG 34
|||||
574 ACATGATGATTTTACGTTGCGTGCAGCGGCTCAATACACAGCGCGGT 525
|||||
34 lThrTrpIleLeuArgSerValArgArgLeuAsnThrAsnArgProArg 50
|||||
524 TTGAAGTCTTCGCGGCTCTTTCATGATGACCGTAGGTCGGCAGCGAG 475
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51 LeuLysSerSerAlaAlaSerLeuMetMetThrValGlySerAlaAlaSe 67
|||||
474 CGGATTTGTCCTTCGTCATTCAGCAAGATGCGAAGCGGCTCGCATCTA 425
|||||
67 rGlyLeuValSerIleAlaLeuThrLysMetAlaAsnGlySerAlaSerT 84
|||||
424 CGGAGGATTTTTCGTAACGCGGCGGTGCGAAGCGCAGTCCATAAGCCT 375
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84 hrAlaGlyIleLeuLeuAsnGlyArgValArgSerAlaValHisLysPro 100  
374 GATTGAATCAGGTTGGCGGCACCTTTTCGCTGCCTCAATTTTCCAGCGC 325  
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117 aSerGly 119

seq\_name: /SIDS1/gcgdata/geneseq/geneseq-emb1/AA2000.DAT:AAV75273

seq\_documentation\_block:

ID AAV75273 standard; Protein; 120 AA.

XX AC AAV75273;

XX DT 21-MAR-2000 (first entry)

XX DE Neisseria meningitidis ORF 628 protein sequence SEQ ID NO:2020.

XX KW Neisseria meningitidis; Neisseria gonorrhoeae; antigen; vaccine;  
XX KW antigenic; diagnosis; immunogenic; infection; meningitis; septicaemia;  
XX KW antibacterial; gene therapy.

XX OS Neisseria meningitidis.

XX XN W09957280-A2.

XX PD 11-NOV-1999.

XX XN 30-APR-1999; 99WO-US09346.

XX PR 01-MAY-1998; 98US-0083758.

XX PR 31-JUL-1998; 98US-0094869.

XX PR 02-SEP-1998; 98US-0098994.

XX PR 02-SEP-1998; 98US-0099062.

XX PR 09-OCT-1998; 98US-0103749.

XX PR 09-OCT-1998; 98US-0103794.

XX PR 09-OCT-1998; 98US-0103796.

XX PR 25-FEB-1999; 99US-0121528.

XX PA (CHIR ) CHIRON CORP.

XX PA (GENO-) INST GENOMIC RES.

XX PI Fraser C, Galeotti C, Grandi G, Hickey E, Masignani V, Mora M;

XX PI Petersen J, Pizza M, Rappuoli R, Ratti G, Scalato E, Scarselli M;

XX PI Tettelin H, Venter JC;

XX DR WPI; 2000-062150/05.

XX DR N-PSDB; AA254035.

XX PT Novel Neisserial polypeptides predicted to be useful antigens for

XX PT vaccines and diagnostics

XX PS Claim 2; Page 1004; 1453pp; English.

XX CC AA253015 to AA254536, AA254577 to AA254615, and AA274253 to AAV75941

XX CC represent novel Neisseria meningitidis and N. gonorrhoeae polynucleotides

XX CC and polypeptides. AA254537 to AA254576 and AA254616 to AA254673 represent

XX CC PCR primers used in the exemplification of the present invention. The

XX CC polypeptides, the polynucleotides, antibodies and compositions of

XX CC the invention can be used as vaccines, as diagnostic reagents, and as

XX CC immunogenic compositions. The polypeptides can be used in the

XX CC manufacture of medicaments for treating or preventing infection due to

XX CC Neisserial bacteria (e.g. meningitis and septicaemia), to detect the

XX CC presence of Neisseria bacteria, or to raise antibodies. They may also

XX CC be used to screen for agonists or antagonists, which may themselves

XX CC have use as antibacterial agents. The polynucleotides of the invention

XX CC may also be used in gene therapy protocols.

XX SQ Sequence 120 AA;

alignment\_scores:

Quality: 532.00 Length: 119

Ratio: 4.626 Gaps: 0

Percent Similarity: 96.639 Percent Identity: 89.916

alignment\_block:

US-09-303-518D-131/rev x AAV75273 ..

Align seg 1/1 to: AAV75273 from: 1 to: 120

674 ATGTGCGTGGCCACTCAAGCGCGGAGGATGGCGCCCAATTCATGTGT 625

|||||  
1 MetCysValProLeuLysProAlaGlyCysGlyProProAsnSerCysVa 17

624 TTCGATATTTGGCAGCATTTTCAGACGGCAGCTCTGGCGCTGCTGCTTAC 575

|||||  
17 lSerMetLeuAlaAlaPheSeraspGlyThrSerAlaProAlaAlaLeuH 34

574 ACATCATGATTTTACGTTTCGCTCAGCGGCTCAATACCAACAGCGCGGT 525

|||||  
34 lThrTrpIleLeuArgSerValLysArgLeuAsnThrSerLysProArg 50

524 TTGAAGTCTTCGCGCGCTCTTTTCATGATGACCGTAGGTCGGCAGCCAG 475

|||||  
51 LeuLysSerSerAlaAlaSerLeuIleThrThrGlySerAlaAlaSe 67

474 CGGATTTGTTCCATCGCATTCAGCAAGATGCGCAACGGCTCGGCATCTA 425

|||||  
67 rGlyLeuValSerIleAlaLeuThrLysMetAlaAsnGlySerAlaSerT 84

424 CGCAGGAGTTTTCGTCGACGGACGGGTGCGAAGCGCAGTCCTCAAGCCT 375

|||||  
84 hrAlaGlyIleLeuLeuAsnGlyArgValArgSerAlaValHisLysPro 100

374 GATTGAATCAGGTTGGCGGCACCTTTTCGCTGCTCAATTTTCCAGCGC 325

|||||  
101 AspTrpIleArgLeuArgArgThrSerSerProLeuLysPheAlaAsnAl 117

324 TTCAGGT 318

|||||  
117 aSerGly 119

seq\_name: /SIDS1/gcgdata/geneseq/geneseq-emb1/AA2000.DAT:AAV82082

seq\_documentation\_block:

ID AAV82082 standard; Protein; 467 AA.

XX AC AAV82082;

XX DT 01-JUN-2000 (first entry)

XX DE Chlamydia pneumoniae antigen CPN100605 protein SEQ ID NO:2.

XX KW Chlamydia pneumoniae; antigen; CPN100605 protein; immunisation;

XX KW vaccine; infection; antibacterial; antiinflammatory; bronchitis;

XX KW community acquired pneumonia; upper respiratory tract infection;

XX KW sinusitis.

XX OS Chlamydia pneumoniae.

XX PN WO200006742-A2.

XX PD 10-FEB-2000.

XX XX 27-JUL-1999; 99WO-IB01331.

XX XX 27-JUL-1998; 98US-0094195.

XX PR 26-JUL-1999; 99US-0361443.

XX PA (CONN-) CONNAUGHT LAB LTD.

PI Murdin AD, Oomen RP;  
 XX WPI; 2000-205466/18.  
 DR N-PSDB; AAZ95378.  
 XX Chlamydia pneumoniae antigens used for immunization and protection  
 PT against Chlamydia diseases  
 XX Claim 6; Fig 1; 48pp; English.

XX The present sequence represents the Chlamydia pneumoniae antigen  
 CC CPN100605 protein. The CPN100605 protein has antibacterial and  
 CC antiinflammatory activities. The Chlamydia pneumoniae CPN100605  
 CC polynucleotide and protein can be used in vaccination methods for  
 CC preventing and treating Chlamydia infection (e.g. infections caused by  
 CC C. trachomatis, C. psittaci, C. pneumoniae or C. pecorum). The  
 CC polynucleotide can be used to produce the protein recombinantly, in the  
 CC construction of vaccine vectors, as a vaccine agent, and in the  
 CC construction of an attenuated Chlamydia strain. The protein are also be  
 CC useful as a vaccine agent, and for the preparation of medicaments for  
 CC treating or preventing Chlamydia infection, e.g. community acquired  
 CC pneumonia, and upper respiratory tract infections such as bronchitis and  
 CC sinusitis.

XX Sequence 467 AA;

alignment\_scores:  
 Quality: 450.50 Length: 470  
 Ratio: 1.532 Gaps: 16  
 Percent Similarity: 62.553 Percent Identity: 29.149

alignment\_block:

US-09-303-518D-131 x AAY82082 ..

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 3 IleThrValAsnArgGlyLeuAspLeuSerLeuGlnGlySerProLysG 19  
 51 GCAAGTCATTTATACGCGCGCGCCATTACCGAAGTCGCGTTCGTCGCG 100  
 19 uSerGlyPheTyrAsn.....LysIleAsp 28  
 101 AAGATATATGTCGC.....ATGCGCCG  
 28 roGluPheValSerIleAspLeuArgProPheGlnProLeuSerLeuLys 44  
 127 ATGAATAATCAAGAGGTGAAGCGGTCAAAAAGCCCAAGTGTCTTTGA 176  
 45 LeuLysValGluGlnGlyAspAlaValCysSerGlyAlaProIleAlaG 61  
 177 AGACAAAAGAAATCGCGCGGTAGTATTACGCGCGCGCTTCAGGCAAA 226  
 61 uTyrLysHisPheProAsnThrTyrIleThrSerHisValSerGlyVal 78  
 227 TCGCGGTATTCACGCGCGCAAGCGGTACTTCAGTCAGTCGTCGATT 276  
 78 alThrAlaIleArgArgGlyAsnLysArgSerLeuLeuAspValIlelle 94  
 277 ...GCCGTTGAAGTCAAGCAAGTTCGAGTTCGAAGCTACGCTACCTGA 323  
 95 LysLysThrProGlyProThrSerThrGluTyr...ThrTyrAspLeuG 110  
 324 AGCGTGGCAAAATGACAGCGCAAAAGTGCAGCGCGCTTCGATTCAT 373  
 110 nThrLeuSerArgSerAspLeuSerGluIlePheLys.....GluA 124  
 374 CAGCGTTATGACGTCGCTCGCACCGCGTTCGTCAGCAAAATCCCTGCC 423  
 124 snGlyLeuPheAlaLeuIleLysGlnArgProPheAsp....IleProAla 139

424 GTAGATGCGGAG...CGTTCCGATCTTCGTCATCGATCGACACAA 470  
 140 IleProThrGlnThrProArgAspValPheIleAsnLeuAlaAspAsnAr 156  
 471 TCCGCTGCTGCCACCGCTACGTCATCATC.....A 502  
 156 gProPheThrProSerProGluLysHisLeuAlaLeuPheSerSerArg 173  
 503 AAGAACGCGCGAAGACTTCAACGCGCGCTGTTGTTGTTGAGCGCGCTG 552  
 173 luGluGlyPheTyrValPheValValGlyValArgAlaIleAlaLysLeu 189  
 553 ACCGAACGTAATTCATGTGTGTAAAGCAGCAGCGCGCAGCGTCGCTC 602  
 206 rGlnGluLeuLysThrIleAlaHisLeuHisThrValSerGlyProPhe 223  
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 603 TCAAAATGCTGCCAATATC...GAACACATCATGAATTTGGCGCGCGCATC 649  
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 650 CTGCGCGCTTGAAGTGGCAGCAGCATTCATTCATCGACGCGCGCGCG 699  
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 700 AATAAA...ACGCTGTGGACCATCAATATCAAGACGTGATGCTATCGG 746  
 240 GluLysGluValValPheThrLeuSerPheGlnAspValLeuThrIleGl 256  
 747 ACGTTTGTTCGTAACAGCGCGCTCGAATACGAGCGCGTGTGCTGCTGG 796  
 256 yHisLeuPheLeuLysGlyArgIleLeuHisGluGlnValThrAlaLeuA 273  
 797 GCGGC.....CTGCAAGTCAAAACCGCGCTCTTCGCGTACCGTTTGG 840  
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 290 GlyAlaSerPheSerSerLeuLeuAsnAspIleSerAspAsnAs 306  
 891 CGCGGTGATTTCGCTGCTGATTGAACGCGTTCGATTCACAAAGCGCGC 940  
 306 pThrLeuIleSerGlyAspProLeuThrGlyArgLeuLysLysLysGluG 323  
 941 ATGAT...TATTGGGACGCTACCAATACAGATTTCGCTTATCGAAGAA 987  
 323 luGluProPheLeuGlyPheArgAspHisSerIleSerValLeuHisAsn 339  
 988 GCGCGCAGCAAGAGCTGTCGCTGCTGCTGCTGCTGCTGCTGCTGCTG 1037  
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 356 oThrPheThrLysThrTyrLeuSerGlyPheLysLysLysArg...T 372  
 1088 AGTTCAG.....ACAGCGCTCAACGCGCGCGCGCGCGCGCGCGCGTA 1128  
 372 hrTyrThrAsnProAspThrAsnLeuHisGlyGluThrArgProIlelle 388  
 1129 CCGATCGGCATTCAGCGCGTAAATGCGTGTGGACATCTCTGCTACCTT 1178  
 389 AspThrAspIleTyrAspLysValMetProMetArgIleProValValPr 405  
 1179 GCTTTTGGCGGATTTAATCGTCGCGCATACCGACAGCGCGCGGCTTTGG 1228  
 405 oLeuLysAlaValIleThrLysAsnPheAspLeuAlaAsnGluLeuG 422  
 1229 GTTCTTGAATTTGACGAGACGCTCGCTTGTGAGCTTCGCTGCTCC 1278  
 422 lPheLeuGluValCysGlyGluAspPheAlaLeuProThrLeuIleAsp 438  
 1279 CCGCGCAATACGAATACGCGCGCTGTTGCGCAAAAGTCTGGAACCAT 1328

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439 ProSerLysThrGluMetLeuThrIleValLysGluSerLeuIleGluTyr 455  
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seq\_name: /SIDS1/gcgdata/geneseq/geneseq-emb1/AA1999.DAT:AAV35375

seq\_documentation\_block:

ID AAV35375 standard; Protein; 469 AA.

XX AAV35375;

XX DT 13-SEP-1999 (first entry)

XX Chlamydia pneumoniae transmembrane protein sequence.

XX Respiratory disease; pneumonia; bronchitis; heart disease; sarcoidosis;  
KW sinusitis; purulent otitis media; erythema nodosum; pharyngitis;  
KW vaccine; neutralising epitope.

XX Chlamydia pneumoniae.

XX PN WO9927105-A2.

XX PD 03-JUN-1999.

XX PF 20-NOV-1998; 98WO-IB01890.

XX PR 04-NOV-1998; 98US-0107078.

XX PR 21-NOV-1997; 97FR-0014673.

XX PA (GEST) GENSET.

XX PI Griffais R;

XX DR WPI; 1999-357842/30.

XX Genome sequence of Chlamydia pneumoniae

XX Page 1170-1171; Disclosure; 1912pp; English.

CC AAV34584-Y35879 represent the proteins encoded by all the open reading  
frames in the complete genome (see AAV34584) of Chlamydia pneumoniae.  
CC C. pneumoniae causes respiratory disease such as pneumonia and  
CC bronchitis and is thought to be a contributing factor in heart  
CC disease, sarcoidosis, sinusitis, purulent otitis media, erythema  
CC nodosum or pharyngitis. The polypeptides encoded by the open reading  
frames of the C. pneumoniae genome (see AAV34584-Y35879) can be used in  
CC immunogenic compositions as vaccines. Vectors containing C. pneumoniae  
CC nucleotide sequences can also be used as immunogenic compositions,  
CC especially where the vector directs the expression of a neutralising  
CC epitope of C. pneumoniae.

XX SQ Sequence 469 AA;

alignment\_scores:

Quality: 450.50 Length: 470  
Ratio: 1.532 Gaps: 16  
Percent Similarity: 62.553 Percent Identity: 29.149

alignment\_block:

US-09-303-518D-131 x AAV35375 ..

Align seg 1/1 to: AAV35375 from: 1 to: 469

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5 IleThrValAsnArgGlyLeuAspLeuSerLeuGlnGlySerProLysG1 21

51 GCAAGTCATTTATGACGGCCCGCCATTACCGAAGTCGCTTGGCG 100

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101 AAGAATATGTCGCC.....ATGCGCGCCC.....TCG 126  
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127 ATGAAATCAAGAGGTAAGCGCTCAAAAAGCCCAAGTGTGTTTGA 176  
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47 LeuLysValGluGlnGlyAspAlaValCysSerGlyAlaProIleAlaG1 63  
177 AGACAAAAGAAATCGCGCGCTAGTATTCTACCGCGCGCTTCAGGCAAAA 226  
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63 uTyrLysHisPheProAsnThrTyrIleThrSerHisValSerGlyVal 80  
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80 alThrAlaIleArgArgGlyAsnLysArgSerLeuLeuAspValIle 96  
277 ...GCCGTTGAGGACGACGAAATCGAGTTCGAACGCTACGTACCTGA 323  
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97 LysLysThrProGlyProThrSerThrGluTyr...ThrTyrAspLeuG1 112  
324 AGCGCTGCAAAATTGAGCAGCGAAAGTGCAGCGCAACCTGATTCAAT 373  
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112 nThrLeuSerArgSerAspLeuSerGluIlePheLys.....GluA 126  
374 CAGGCTTATGACTCGCGTTCGACCGCTCGTTTCAGCAAAATCCCTGCC 423  
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126 snGlyLeuPheAlaLeuIleLysGlnArgProPheAsp...IleProAla 141  
424 GTAGATGCCGAG...CGTTCCGCTCTTCGTCAATCGGATGGACACCA 470  
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142 IleProThrGlnThrProArgAspValPheIleAsnLeuAlaAspAsnAr 158  
471 TCCGCTGGCTGCCGACCTACGGTCATCATC.....A 502  
|||:||||: |||  
158 gProPheThrProSerProGluLysHisLeuAlaLeuPheSerSerArg 175  
503 AAGAAGCGCGCAAGACTTCAACCGCGCTGTGTTGTTATGAGCGCGCTG 552  
|||:||||: |||  
175 luGluGlyPheTyrValPheValValGlyValArgAlaIleAlaLysLeu 191  
553 ACCGAACGTAAATCCATGTGTGTAAAGCAGCAGCGCGCACACGTGCGTC 602  
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192 PheGlyLeuArgProHisIleValPheArgAspArgLeuThrLeuProth 208  
603 TGAATATGCTGCCAATATC...GAACACATGAATTTGGCGCGCGCATC 649  
|||:||||: |||  
208 rGlnGluLeuLysThrIleAlaHisLeuHisThrValSerGlyProPhe 225  
650 CTGCGCGCTTACGTGCGCACCATTCATTTCATCGAGCCAGTCGGCGCG 699  
|||:||||: |||  
225 roSerGlySerProSerIleHisIleHisSerValAlaProIleThrAsn 241  
700 AATAAA...ACCGTGTGACCATCAATTATCAAGACGTGATTGCTATCGG 746  
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242 GluLysGluValValPheThrLeuSerPheGlnAspValLeuThrIleG1 258  
747 ACCTTTCTGTAACAGCGCTCTGAATACCGCGCGCTGTTGCTGCTGG 796  
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258 yHisLeuPheLeuLysGlyArgIleLeuHisGluGlnValThrAlaLeuA 275  
797 GCGGC.....CTGCAAGTCAACAAACCGCGCTCTTGGCTACCGCTTTG 840  
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275 laGlyThrAlaLeuLysSerSerLeuArgArgIleThrThrLys 291  
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292 GlyAlaSerPheSerSerLeuIleAsnLeuAsnAspIleSerAspAsnAs 308  
891 CCGCGTCAATTCGGTTCGGTATTGAACGTCGGATTGCACAGGCGCGC 940  
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179 heValValGlyValGlnAlaAlaLeuPheGlyLeuLysProHis 195
179  |||... ..|||
571 GTGTGTAAACGACGCGGAGCGGTCTGTGAATAATGCTGCCAATAT 620
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196 IleIleSerThrAspArgLeuThrLeuProThrGlnAspLeuValSerIle 212
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212 eAlaHisLeuHisThrIleAspGlyProPheProSerGlySerProSerT 229
212  |||... ..|||
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668  |||... ..|||
229 hrHisIleHisIleAlaAraGleArgAsnGluArgAspValValPhe 245
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715 ACCATCAATTAACAGACGTGATGTCATCGGACGTTGTCGTACACAGG 764
715  |||... ..|||
246 ThrIleSerPheGlnGluValLeuSerIleGlyHisLeuPheLeuLysG 262
246  |||... ..|||
765 CGCTCTGAATACCGAGCGGTGTTGCCCTTGGCGGC...CTGCAAG 808
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262 yPheValLeuGlyGlnGlnIleValAlaLeuAlaGlySerAlaLeuProp 279
262  |||... ..|||
809 TCACAAACCGCGCTCTTCGTCACGTTTGGTGGCGAAGTGTCACAA 858
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859 CTTACCGCGCGGTGTTGTCGCGGACAAAC...CGCGTCAATTCGCG 905
859  |||... ..|||
296 LeuLeuProLysAspIlePheSerSerAspGluIleThrLeuIleSerG 312
296  |||... ..|||
906 TTCGGTATGACGGTGGCATTCACAGCGCGCATGAT...TATTTGG 952
906  |||... ..|||
312 yAspProLeuThrGlyArgLeuCysLysLysGluGluAsnProCysLeuG 329
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953 GACGCTACCAACATCAGATTCCTGATTCGAGAGAGCGCGCAGCAAGAG 1002
953  |||... ..|||
329 lyMetArgAspHisThrIleThrLeuProAsnProLysThrArgGlu 345
329  |||... ..|||
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1003  |||... ..|||
346 SerPheSerPheLeuArgLeuGlyTrpAsnLysLeuThrValThrArgTh 362
1053 CACTCTCGCGCATTCCTTAAACAA...CTCTCAAGTTCACGA 1096
362  |||... ..|||
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379 hrAsnMetHisGlyGluLysArgProIleIleAspAlaGluIleThrGlu 395
1147 CGCGTAATCGCGTGGACATCGCTGCTACCTGCTTTTTCGCGGATTTAAT 1196
1147  |||... ..|||
396 ArgValSerAlaIleProValProValAlaLeuIleIleLysAlaLeuG 412
1197 CGTCGCGATACCGACGCGCGCGGTGTTGGTGTGTTGGAATGGACG 1246
412  |||... ..|||
412 uThrGlnAsnPheGluGluAlaCysArgLeuLeuLeuGluValAlaP 429
1247 AAGAGACCTCGCTTGTGACGTCGTCGCGCGGCAATACGAAATAC 1296
429  |||... ..|||
429 roGluAspPheAlaLeuProThrPheIleAspProSerLysThrGluMet 445
1297 GGCGCGTGTTCGCAAGTCTCGAACCATTGAGAG 1335
446  |||... ..|||
446 PheSerIleValLysGluSerLeuLeuArgThrGlnLys 458
seq_name: /SIDSI/gcgdata/geneseq/geneseq-emb1/AA1999.DAT:AA1999
seq_documentation_block:
ID AAY34467 standard; Protein; 443 AA.
XX
AC AAY34467;

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XX
DT 25-AUG-1999 (first entry)
XX
DE Porphyromonas gingivalis protein PG122.
XX
KW Porphyromonas gingivalis; PG: periodontal disease; gingivitis;
XX vaccine; antigenic.
XX
OS Porphyromonas gingivalis.
XX
PN WO9929870-A1.
XX
PD 17-JUN-1999.
XX
PF 10-DEC-1998; 98WO-AU01023.
XX
PR 04-AUG-1998; 98AU-0005028.
PR 10-DEC-1997; 97AU-0000839.
PR 31-DEC-1997; 97AU-0001182.
PR 30-JAN-1998; 98AU-0001546.
PR 10-MAR-1998; 98AU-0002264.
PR 09-APR-1998; 98AU-0002911.
PR 23-APR-1998; 98AU-0003128.
PR 05-MAY-1998; 98AU-0003338.
PR 22-MAY-1998; 98AU-0003654.
PR 29-JUL-1998; 98AU-0004917.
XX
(CSLC-) CSL LTD.
XX
PA Agius CH, Barr IG, Hocking DM, Margetts MB, Patterson MA;
PI Ross BC, Rothel LJ, Webb EA;
XX
DR WPI; 1999-385613/32.
DR N-PSDB; AAX91685.
XX
PT Antigenic Porphyromonas gingivalis peptides for preventing
PT gingivitis
XX
PS Claim 1; Page 445-446; 588pp; English.
XX
CC AAX91536 to AAX91801 encode two hundred and sixty six antigenic
CC Porphyromonas gingivalis (PG) polypeptide sequences given in AAY34318 to
CC AAX34583. AAX91802 to AAX91989 represent PCR primers used in the
CC isolation of the PG polypeptides. The PG polypeptides have antibacterial
CC activity with a vaccine mechanism of action. The PG polypeptides can be
CC used as vaccines especially against Porphyromonas gingivalis. Probes can
CC be used to detect Porphyromonas gingivalis in standard hybridisation
CC assays. Porphyromonas gingivalis is involved in periodontal disease
CC especially gingivitis.
XX
SQ Sequence 443 AA;

```

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alignment_scores:
  Quality: 161.50      Length: 490
  Ratio: 0.699        Gaps: 22
  Percent Similarity: 47.143  Percent Identity: 20.612

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alignment_block:
US-09-303-518D-131 x AAY34467 ..

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Align seg 1/1 to: AAY34467 from: 1 to: 443
```

```

37 GCGGCGACGCGGAGCAAGTCATTTATGACGCGCGCCATACCGAAGT 86
|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||
19 AlaGlyLysProValGluValLeu.....ProIleProSerGlnVa 32
|||||:|||||:|||||:|||||:|||||:|||||:|||||
87 CGCGTTGCTGCGAAGAATAATGTCGCGCATGCGCCCTCGATGAAATCA 136
|||||:|||||:|||||:|||||:|||||:|||||:|||||
32 lValIleProLeuGlyGlnHisIleGlyAlaProAlaThrVal 49
|||||:|||||:|||||:|||||:|||||:|||||:|||||
137 AGAAGGTGAAGCGGTCAAAAAAGCCCAAGTCTGTTTGAAGACAAAAG 186
|||||:|||||:|||||:|||||:|||||:|||||:|||||

```



49 yLysGlyAspGluValLysValGlyThrIleIle ..... 60  
187 AATCCGGCGGTAGTATTACTGCGCGGCTTCAGCAAAATCCCGCT.. 234  
61 .....AlaGlnAlaGlyGlyPheValSerAlaAs 70  
235 .ATTCACCGTGGC...GAAAGCGGTACTTCAG..... 264  
70 nIleHisSerValSerGlyLysValLysValLysIleAspAsnValTyrA 87  
265 .....TCAGTCGTGATTGCGGTTGAAGCAAC 291  
87 spSerSerGlyTyrProLysProAlaValPheIleSerValGluGlyAsp 103  
292 GAGCAATCGAGTTCGACCGCTACGTA..... 318  
104 .....GluTrpGluGluGlyIleAspArgSerProAlaIleVally 117  
319 .....CCTGAGCGCTGGCAAAATTCAGCAGCGAAA 349  
117 sGluCysAsnLeuAspAlaLysGluIleValAlaIleSerAlaAlaG 134  
350 AAGTCGCGCGCAACCTGATTCATCAATCAGCGTTCGACTGCGCTTCGCACC 399  
134 lyIle.....ValGlyLeuGlyGlyAla.....Thr 142  
400 CGTCCG.....TTCCAGCAAAATCCCTGCGGTAGATCGCGAGCC 437  
143 PheProThrHisValLysLeuSerProProGlyAsnLysAlaGlu.. 158  
438 GTTCGCGCATCTTCGTCATCGATGGACACCAATCCGCTGCGCGCAGCC 487  
159 .....IleLeuIleIleAsnAlaValGluCysGluProTyrLeuThrSerA 174  
488 CTACGGTCATCATCAAGAGCCGCCGAGACCTCAACCGCGGCTGTG 537  
174 spHisValLeuMetLeuGluHisGlyGluIleMetIleGlyValSer 190  
538 GTATTGAGCGCGCTGACCGAGCGTAAATCCATGCTGTGTAAGCAGCA.. 585  
191 IleLeuMet.....LysAlaIleGlnValAsnLysAlaValI 203  
586 .GGCGCAGACGTGCGCTGTGAAATGCT..... 612  
203 eGlyValGluAsnAsnLysLysAspAlaIleAlaHisLeuThrLysLeuA 220  
613 .....GCCAATATCAACACATGAATTTGGCGCGCCGCACTCCT 651  
220 laThrAlaTyrProGlyIleGluValMetProLeuLysValGlnTyrPro 236  
652 GCGCGCTTGAGTGGCAGCACCATTCATTCATC..... 684  
237 GlnGlyGlyGluLysGlnLeuIleAspAlaValIleArgLysGlnVally 253  
685 .....GAGCAGTCGGCGGCAATAAACCGTGTGGACCATCAATT 724  
253 sSerGlyAlaLeuProIleSerThrGlyAlaValValGln.....AsnV 268  
725 ATCAAGACGTGATTCGTCGACGCTTTGTCGTAACAGCGCTCTGAAT 774  
268 aGlyThrValPheAlaValTyrGluAlaValGlnLysAsnLysProLeu 284  
775 ACCGAGCGCGTGTGCTTCGCGCGGCTGCAAGTCAACAAACCGCGCT 824  
285 ValGluArgIleValThrValThrGlyLysLysLeuSerArgProSerAs 301  
825 CTTGGTACCGTTTGGGTGGCGAAGGTCTCAACTT.....ACCGCG 868  
301 nLeuLeuValArgIleGlyThrProIleAlaAlaLeuIleGluAlaAlaG 318  
869 GCGAATTTGTTGTCGCGGACAAACCGGTGATTTCGTTTCGTTGTAAC 918  
318 lyGlyLeuProGluAsnThrGlyLysIleIleGlyGly..... 331

919 GGTCCGATTGCACAGCGCGCCATGATTATTTGGACGCTACCAACAATCA 968  
331 ..... 331  
969 GATTTCCTGTTATCGAAGAAGCGCGCAGCAAAAGAGCTGTTCGGCTGGTTG 1018  
332 .....ProMetMetGlyArgAlaL 338  
1019 CCCCACCGCGGACAAATACTCCATCCAGCGCACCACCTCTCGGCCAATTC 1068  
338 euLeuSerProAsp...ValProValThrLysGlySerSerGlyValLeu 353  
1069 CTAAAAAAACAAACTCTTCAAGTTCACGACAGCGCTCAACAGCGCGGACCG 1118  
354 lleLeuAspArg.....GluGluAlaValArgLysProMetAr 366  
1119 CCCCATGTACGATCGGCACCTTATGACGCGGTATGCGCTTGGACATCC 1168  
366 gAspCysIleArgCysAlaLysCysValGlyValCysProMetGlyLeuA 383  
1169 TGCCTACCTGCTTTCGCGGATTTAATCGTCGCGGATACCGACAGCGCG 1218  
383 snProAlaPheLeuMetArgAspThrLeuTyrLysSerTrpGluThrAla 399  
1219 CAG.....GCTTTGGGTGCTTGGAAATTTGGACGAGGAAGACCTCGC 1259  
400 GluLysGlyAsnValValAspCysIleGluCysGlySer..... 412  
1260 TTTGTGACGCTTCGTCCTCGCGGCAATACCAATACGCGCGCTGTGTC 1309  
413 ....CysSerPheThrCysProAlaAsnArgProLeuLeuAspTyrIleA 428  
1310 GCAAGTGTCTGGAACCACTT 1329  
428 rgGlnAlaLysIleThrVal 434  
seq\_name: /SIDSL/gcgdata/geneseq/geneseq-emb1/AA1999.DAT:AA1999

seq\_documentation\_block:  
ID\_AAY34343 standard; Protein: 451 AA.  
AC AAY34343;  
DT 25-AUG-1999 (first entry)  
DE Porphyromonas gingivalis protein PG122.  
KW Porphyromonas gingivalis; PG; periodontal disease; gingivitis;  
KW vaccine; antigenic.  
OS Porphyromonas gingivalis.  
PN WO9929870-A1.  
PD 17-JUN-1999.  
PF 10-DEC-1998; 98WO-AU01023.  
XX 04-AUG-1998; 98AU-0005028.  
PR 10-DEC-1997; 97AU-0000839.  
PR 31-DEC-1997; 97AU-0001182.  
PR 30-JAN-1998; 98AU-0001546.  
PR 10-MAR-1998; 98AU-0002264.  
PR 09-APR-1998; 98AU-0002911.  
PR 23-APR-1998; 98AU-0003128.  
PR 05-MAY-1998; 98AU-0003338.  
PR 22-MAY-1998; 98AU-0003654.  
PR 29-JUL-1998; 98AU-0004917.  
XX (CSLC-) CSL LTD.  
XX Agius CT, Barr IG, Hocking DM, Margetts MB, Patterson MA;  
PI





```

alignment_scores:
  Quality: 145.50      Length: 497
  Ratio: 0.677        Gaps: 27
  Percent Similarity: 43.260  Percent Identity: 24.748

alignment_block:
  US-09-303-518D-I31 x AAB59826  ..

Align seg 1/1 to: AAB59826 from: 1 to: 1615

38  CGGSCAGACGGGACGAGTCATTATGACGGCCGCCATTACCGAGTC 87
||||| : : : : : : : : : : : : : : : : : : : : :
699  ArgAlaSerSerProLysSerIleSerProLysProArgProThrCy 715
||||| : : : : : : : : : : : : : : : : : : : : :
88  GCCT.....TGCTGGCGAAGATATGTGGGATGCGCCCTCGAT 128
||||| : : : : : : : : : : : : : : : : : : : : :
715  sArgProSerProGlyThrAlaArgArgValSerThrThrSerProArg. 731
||||| : : : : : : : : : : : : : : : : : : : : :
129  GAAATCAAGGAGGTGAACCGCTCAAAAAGCCCAAGTCTCTTTGAAG 178
||||| : : : : : : : : : : : : : : : : : : : : :

```

```

alignment_scores:
  Quality: 145.50      Length: 497
  Ratio: 0.677        Gaps: 27
  Percent Similarity: 43.260  Percent Identity: 24.748

alignment_block:
  US-09-303-518D-131 x AAB59826  ..

  Align seg 1/1 to: AAB59826 from: 1 to: 1615

38  CGGCGACAGCGGACCAAGTCATTATGACAGCGCCGGCCATTACCGAAGTC 477
|||||:  |||  ::  ::  ::  ::  ::  ::  ::  ::  ::  ::  ::  ::  |||
699  ArgAlaSerSerProLysSerIleSerProLysProArgProThrCy 715
|||||:  |||  ::  ::  ::  ::  ::  ::  ::  ::  ::  ::  ::  ::  |||

88  GCGT.....TGCTTGCGAAGATAATGTCGGCATGCGCCCTCGAT 128
|||  |||  |||  |||  |||  |||  |||  |||  |||  |||  |||  |||
715  sArgProSerProGlyThrAlaArgValSerThrThrSerProArg. 731
|||||:  |||  ::  ::  ::  ::  ::  ::  ::  ::  ::  ::  ::  ::  |||

129  GAAATCAAGGAAGGTGAAGCCGTCACAAAAGCGCAAGTGCTCTTTGAAG 178
|||||:  |||  |||  |||  |||  |||  |||  |||  |||  |||  |||  |||
732  SerThrThrGlyArg.....ArgTtpSerSerProAla..... 742
|||||:  |||  ::  ::  ::  ::  ::  ::  ::  ::  ::  ::  ::  ::  |||

179  ACAAAGATATCGCGGTAGTATTACTCGCGCGGCTCAGGCAAAATC 228
|||||:  |||  |||  |||  |||  |||  |||  |||  |||  |||  |||  |||
743  .ArgArgSerAlaGlyArgAlaGlyArgAlaGlyCysAlaArgSerSerA 759
|||||:  |||  ::  ::  ::  ::  ::  ::  ::  ::  ::  ::  ::  ::  |||

229  GCGCGATTCA...CGGTGCGGAAAAGCGGTACTTCAGTCAGTCGTGAT 275
|||||:  |||  |||  |||  |||  |||  |||  |||  |||  |||  |||  |||
759  rGlyThrSerArgProIleArgSerAlaArgProSerCysSerLysSer 775
|||||:  |||  ::  ::  ::  ::  ::  ::  ::  ::  ::  ::  ::  ::  |||

276  TGCGGTGTGAAGCAACGACGAATCGAGTTCGACGCTACGTACCTGAAG 325
|||||:  |||  ::  ::  ::  ::  ::  ::  ::  ::  ::  ::  ::  ::  |||
776  .ProThrSerValSer.....AlaPheProProS 785
|||||:  |||  ::  ::  ::  ::  ::  ::  ::  ::  ::  ::  ::  ::  |||

326  CGCTGCAAAATTGACGCGCAAAAAGTCGCGCAACCTGATTCAATCA 375
|||||:  |||  |||  |||  |||  |||  |||  |||  |||  |||  |||  |||
785  eProAlaArgAlaSerArgThrArgCysArgArgAsnSerLeuProSer 801
|||||:  |||  ::  ::  ::  ::  ::  ::  ::  ::  ::  ::  ::  ::  |||

376  GGCTTAGGACTCGCTTCGACCCCT.....CCGTTTCAGCAAAAT 416
|||||:  |||  |||  |||  |||  |||  |||  |||  |||  |||  |||  |||
802  SerValThrArgSerSerAlaThrArgAlaAlaThrProArgArgLysTh 818
|||||:  |||  ::  ::  ::  ::  ::  ::  ::  ::  ::  ::  ::  ::  |||

417  CCCTGCGGTAGATGCGGACCGGTCGCGCATCTCGTCA..... 454
|||||:  |||  ::  ::  ::  |||  |||  |||  |||  |||  |||  |||  |||
818  rProCysCysGlyArg.ThrThrArgProProSerSerThrArgAsnSer 834
|||||:  |||  ::  ::  ::  ::  ::  ::  ::  ::  ::  ::  ::  ::  |||

455  .....ATGCGGTGACACCAATCCGCTGG.....CT 480
|||||:  |||  |||  |||  |||  |||  |||  |||  |||  |||  |||  |||
835  SerArgAlaThrTnpMetArgTrpAsnSerSerArgTrpAsnValArgph 851
|||||:  |||  ::  ::  ::  ::  ::  ::  ::  ::  ::  ::  ::  ::  |||

481  GCGGACCTACGTCATCATCAAGAGCGCGCGAAGACTTCAACACGGG 530
|||||:  |||  |||  |||  |||  |||  |||  |||  |||  |||  |||  |||
851  eProSerMetAlaProAlaSerArgAlaAlaProThrAlaLysSerArg 868
|||||:  |||  ::  ::  ::  ::  ::  ::  ::  ::  ::  ::  ::  ::  |||

```

```
531 CCTGTTGGTATTGAGCGCCTGACCGCAACGTAATAATCCATGTTGTTAAAG 580
  ::::::::::: ||| ::::
868 lyArgThrIleCysSerSerPro.....SerAlaAlaProThr 881
  ::::::::::: ||| ::::
581 CAGCAGCGCAGAGCTGCGCTCTGAAATGCTGCAATATCGAAACACAT 630
  ::::::::::: ||| ::::
882 ProArgAlaArgThrProAlaThrProThrProSerSerArgGlnPr 898
  ::::::::::: ||| ::::
631 GAATTTGGCGGCGCGC.....ATCCTGCGCGCTT 659
  ::::::::::: ||| ::::
898 oSerGlySerAlaArgProSerProProSerSerSerAlaIleProArg 915
  ::::::::::: ||| ::::
660 GAGTGGCAGCCACA.....TTCATTTCATCGAGC..... 688
  ::::::::::: ||| ::::
915 rgThrAlaAlaArgArgCysAlaGlyPheSerSerAlaAlaThrAsp 931
  ::::::::::: ||| ::::
689 .....CAGTGGCGCGCAATAAACCGTG 711
  ::::::::::: ||| ::::
932 SerAlaIleArgArgSerSerThrThrArgSerAlaArgSerArgAs 948
  ::::::::::: ||| ::::
712 TGGACCATCAATTATCAAGAGCTGATTCGCTATCGAGCTTGTTCGTAAC 761
  ::::::::::: ||| ::::
948 nThrProSerSerAlaSerThr..... 955
  ::::::::::: ||| ::::
762 AGCGCTCTGAATACCGAGCGCTGGTTCCTTGGCGGCGCTGCAAGTCA 811
  ::::::::::: ||| ::::
956 .....AlaThrAlaPro.....ProThrArgLysPro 964
  ::::::::::: ||| ::::
812 ACAAACCGCGCTCTGCGTACCCTTTTGGGTGCGAAGGTCTCTCAACTT 861
  ::::::::::: ||| ::::
965 ThrThrGlySerThrCys.....CysAlaCysAr 974
  ::::::::::: ||| ::::
862 ACCGCGCGCAATTGGTTCAGCGGACACCGCGTGATTTCCGGTTCGGT 911
  ::::::::::: ||| ::::
974 gProAlaSerThrValAlaAlaArgArgLys.....ProValArg. 988
  ::::::::::: ||| ::::
912 ATTGAACGGTGGATTCACAAAGCGCGCATGATTATTGG..... 952
  ::::::::::: ||| ::::
989 .....LysValAlaAlaGlnSerSerArgProSerCysTrpLysSerArg 1003
  ::::::::::: ||| ::::
993 .....GACGTACCAATCAGATTTCCTTATCGAAGAGCGCGCAGC 996
  ::::::::::: ||| ::::
1004 SerMetThrAlaThrThrGlyArgThrProThrCysAsnSerAlaArgAr 1020
  ::::::::::: ||| ::::
997 AAAGAGCTGTTCGGTGGTTCGCGCGCAGCGGACAAATCTCATCAC 1046
  ::::::::::: ||| ::::
1020 gProValIleSer.....ArgArgSer.....ProSerA, 1030
  ::::::::::: ||| ::::
1047 GC.....GCACCACTCTCGGCAATTTCCTAAACAACTCTTCAAGT 1090
  ::::::::::: ||| ::::
1030 rgMetPheGlyArgLeuSerAlaSerIleAsnMetArgSerThrSer 1046
  ::::::::::: ||| ::::
1091 TCACGACAGCCCTCAACGCGCGCGACCGCGCATCGGTACCGATCGGCAC 1140
  ::::::::::: ||| ::::
1047 ValSerAlaProArgThrCys.....ArgAlaThrSe 1057
  ::::::::::: ||| ::::
1141 TATGACCGGTAATGCGGTGGACATCTGC..... 1171
  ::::::::::: ||| ::::
1057 rSerSerAlaSerCysA-gCysLeuSerCysProGlnSerThrThrAlaA 1074
  ::::::::::: ||| ::::
1172 .....CTACCTTCCTTTTGGCGGATTAAATCGTCGCGC 1204
  ::::::::::: ||| ::::
1074 laTrpAsnSerGlyTrpThrProAlaProCysProSerProMetAla 1090
  ::::::::::: ||| ::::
1205 ATACCGACAGCGCGCAGG.....CTTTGGTTCG 1233
  ::::::::::: ||| ::::
1091 GlyThrThrArgSerArgSerSerArgArgThrProSerTrpProSe 1107
  ::::::::::: ||| ::::
1234 TTGGAATTCGACGAGAACACCTCGCTTTGTGCAGCTTCGCTGCCCGGG 1283
  ::::::::::: ||| ::::
1107 rArgAsnTrpTySerArgArgArgAsnThrProSerSerAsnSerAlaL 1124
  ::::::::::: ||| ::::
```

```
132 AATCAAGGAAGTGAAGCCG..... 151
   :::::
650 spAlaArgAsnArgAspProLeuGlyArgGlyCysThrAspProCysCys 666
152 .....TCAAAAAGCCCAAGTCGCTTTGAAGACAAAGA... 187
   :::::
667 ProProAlaThrArgArgGlyArgAsnCysCysArgAlaAlaArgAr 683
188 .....ATCCGGGGCTAGTATTACTGCGCGGCTTCAGGCAA 224
   :::::
683 gPheArgGlyProLeuArgAlaAlaProAlaCysAlaArgPheAla 699
225 AATCCCGCTATTACCG..... 242
   :::::
700 TyrArgArgLeuGlyProArgCysThrSerArgGlyArgSerArgCysSe 716
243 .....TGCGAAAGCCGCTACTTCAGTCAGTCGTG 273
716 rProAspArgCysCysArgTrpSerArgCysSerProSerProArgA 733
274 ATTGC..... 278
733 rGysProProSerSerProAlaGlyAlaProGlyAlaThrCysSerArg 749
279 .....CGTTGAAGCAA.....CGACGAATCGAGT 304
750 ArgProPheSerArgSerArgAspSerAlaGlyProArgAlaAlaArgph 766
305 TCGAAGCGTAGTACCTGAACCGCTGGCAAAATTGACGAGCGAAAAGTG 354
766 eArgArgCysArg.....AspAlaCysGluArgArgAlaArgC 779
355 CGCGCAACCTGATTCAATCAGGCTTATGGAAGTTCGCGCCCGCTCC 404
779 ysProGlyPro.....ArgSerAlaProSer 787
405 GTTCAGCAAAATCCTCGCTAGATCGGAGCGCTTCGC..... 443
788 IleArgArgGlySerArgAspArgSerArgAlaSerArgSerArgGlySe 804
444 .....CATCTTCGTCATGCGC 459
804 rProLeuCysGlyAlaThrAlaThrSerCysProArgArgArgCysS 821
460 ATGGACACCAATCGCTGGCTGC.....CGACCC 488
821 erIleGlyAlaSerSerGlyCysProHisProProValaArgSerPro 837
489 TACGTCATCATCAAGAAGCCGCGAGACTTCAACGCGGCTGTGG 538
838 ValAsnSerSerLysArgAlaHisArgCysThrAlaArgArgGlyAr 854
539 TATTGACCCGCT.....GACCGAAGCTAAATCCATGTGTGATAA 579
854 gPheArgGlyProThrSerArgAspThrGlyArgArgCysTrpArgT 871
580 GCACGAGCGGACGCTGCGCTGAAATGCTGCGCAATATCGAACACA 629
871 rpProArgProArgArgCys.....ArgCysSerArgArgTrpGlyArg 885
630 TGAATTGGCGCGCCGCTGCTGCGGCTGAGTGCACCGCCACATTCATT 679
886 ProLeuTrpAlaSerGlyCysProArgAlaArgTrp..... 897
680 TCATCGAGCCGAGTCGGCGGCAATAAACCGCTGTGACCATCAATTATCAA 729
898 .ArgArgGlySerAsnTrpSerSerGlyArgSerSerAlaAlaSerProL 914
730 GAGGTGATTGCTATCGACGCTTTGTCGTAACAGCGGCTCTGAATACCGA 779
914 ysArgThrCysGlyArgArg...ValArgSerAspThrSerAlaArgArg 929
780 GCGCGT..... 785
```

```
930 SerArgCysProAlaSerSerProIleArgTrpThrGlyArgCysArgAr 946
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786 .....GGTTGCCCTTGGCGGCTCGCAAGTCAACAAACCG 819
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946 gTrpArgArgProLeuGlyCysSerProArg.....A 957
820 CGCCTCTTGCTACCGTTTGGTGGGAGGTTCTCAACTTAC..... 863
   :::::
957 laThrCysThrAlaArgCysGlyArgAspGlyCysSerAlaPheGly 973
863 ..... 863
974 AsnProLeuHisArgSerLeuArgGlyProTrpAlaAlaProPheArgAl 990
864 .....CGCGCGCAATTGTTGACGCGCACACCCG 894
   :::::
990 aHisArgSerArgSerThrThrArgArgCysAlaValArgGlySerSerA 1007
895 GTGATTTCGGTTCGGTATTGAACGGTGCATTGCACAAG..... 934
   :::::
1007 rHisAspArgThrAlaSerThrArg..ArgProHisLysProProLysG 1023
935 .....GCGCGCATGATTATTGGGACGCTACCACA 964
1023 lyCysAlaThrAspIleHisSerGlyArgTyrcysTrpProArgThrAla 1039
965 ATCAGATTTCGGTTATCGAAGAGCGCGCAGCAAGAGCTGTTCGGCTGG 1014
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1040 .....SerSerArgAlaAlaSerGlyAlaSerAlaLysAr 1051
1015 GTTGCAGCGCGCAGCGACAAATCTCCATCAGCGCCACCA..... 1054
1051 gThrArgLeuArgArgArgSerCysProValArgSerProArgArgArgG 1068
1055 ..CTCTCGGCCATTTCCTAAAAACAACACTTCAAGTTTCAGCAGACGCG 1102
   :::::
1068 lyThrArgAlaAlaTrpHisSerAlaCysGlySerSerSerArgArgPro 1084
1103 TCNACGCGCGCGACCGCGCATGG.....TACCGATCGGCATTTAT 1143
1085 SerSerGly.....ArgProTrpSerValProIleArgProSerSerIl 1099
1144 GAGCGCGTAAATCGCGTTGCA..... 1163
1099 eCysGly.ArgAlaValGlyLeuThrSerProSerSerProLeuAsnArg 1115
1164 .....CATCCTGCCTACCTTGTCTTTTGGCGGATTTAA 1195
1116 ProPheAlaArgArgSerAlaProAlaSerThrProCysArgArgHisA 1132
1196 TCGTCGCGGATACCGACGACGCGAGGCTTGGTGTGCTTGAATTGGAC 1245
1132 nArgArgArgTyrglySerArgArgProPhe.....A 1143
1246 GAAGAAGA 1253
1143 rArgArg 1145
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seq\_name: /SIDS1/gcgdata/geneseq/geneseq-emb1/AA2001.DAT:AA859817  
seq\_documentation\_block:  
ID AA859817 standard; Protein; 999 AA.  
XX AA859817;  
XX  
XX  
DT 04-APR-2001 (first entry)  
XX  
DE TutD protein #8.  
XX  
KW Toluene degradation; enzyme; waste degradation; TutD.  
XX  
OS Thauera aromatica.

OS Xanthomonas maltophilia.  
 OS Geobacter metallireducens.  
 XX Azarcus toluolyticus.  
 PN WO200072650-A2.  
 XX 07-DEC-2000.  
 XX  
 PF 24-MAY-2000; 2000WO-US14298.  
 XX  
 PR 01-JUN-1999; 99US-0323872.  
 XX  
 PA (UYOH-) UNIV OHIO.  
 XX  
 XI Coschigano PW;  
 XX  
 XX WPI; 2001-041080/05.  
 DR N-PSDB; AAF23625, AAF23627.  
 XX  
 PT Composition comprising toluene degrading enzyme useful for biological  
 PT treatment of organic compounds, especially for degrading toluene or its  
 PT analogs -  
 XX  
 PS Disclosure; Fig 5; 122pp; English.  
 XX  
 CC The present invention relates to toluene degrading enzyme genes and  
 CC proteins tutH (see AAF23629 and AAB59831), tutI (AAF23630 and AAB59832),  
 CC tutF (AAF23631 and AAB59833) and tutG (AAF23632 and AAB59834). The  
 CC toluene degrading enzymes are homologues of pyruvate formate lyase. The  
 CC toluene degrading enzymes are useful for biological treatment of organic  
 CC compounds and in particular for the degradation of toluene and its  
 CC analogs contained in liquid or solid waste source. The present sequence  
 CC is a protein sequence for toluene degrading enzyme, TutD.  
 XX  
 XX Sequence 999 AA;

alignment\_scores:  
 Quality: 129.00 Length: 573  
 Ratio: 0.617 Gaps: 27  
 Percent Similarity: 36.475 Percent Identity: 22.339

alignment\_block:

US-09-303-518D-131 x AAB59817 ..

Align seg 1/1 to: AAB59817 from: 1 to: 999

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32 CCATCGCGGGGAGCGGAGCAAGTCAATTAATGACGGCGCGCCATTACC 81
||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||
25 ProSerGlyAlaSerArgSerSerAlaAlaAlaProArgArgProPr 41
82 GAAGTCGGTTCCTGGCGAAGATATGCGCATGCGCCCTCGATGAA 131
|::: ||||| ||||| ||||| ||||| ||||| ||||| |||||
41 oSerHisSerSerAlaAlaTyrGlyAlaSerCysThr.....A.55

132 AATCAAGGAGGTGAAGCG..... 151
:::||||| |||||
55 spAlaArgAsnArgAspProLeuGlyArgGlyCysThrAspProCys 71
152 .....TCAAAAAGCCCAAGTCTGTTTGAAGACAAAGA... 187
72 ProProAlaThrArgArgLysArgAsnCysCysArgAlaAlaArg 88
188 .....ATCCGGCGGTAGTATTACTGCGCGCGCTTCAGGCAA 224
:::||||| ||||| ||||| ||||| ||||| ||||| |||||
88 gpheargGlyProLeuArgAlaAlaProAlaCysAlaArgPheAla 104
225 AATCGCCGCTATTCAACG..... 242
||||| |||||
105 TyrArgArgLeuGlyProArgCysThrSerArgGlyArgSerArgCys 121
243 .....TGCGGAAAGCGCGTACTTCACTCAGTCGTCGT 273
||| ||||| ||||| ||||| ||||| ||||| |||||

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121 rProAspArgCysCysArgTrpSerArgCysSerSerProSerProArg 138
274 ATTGC..... 278
138 rgCysProProSerSerProAlaGlyAlaProGlyAlaThrCysSerArg 154
279 .....CGTTGAAGCAA.....CGACGAAATTCGAGT 304
155 ArgProPheSerArgSerArgAspSerAlaGlyProArgAlaAlaArgph 171
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355 CGCGCGCAACCTGATTCAATCAGGCTTATGGACTTGGCTTCGCACCGCTCC 404
184 ysProGlyPro.....ArgSerAlaProSer 192
405 GTTCAGCAAAATCCCTGCGGTAGATGCCGAGCGCTTCGC..... 443
193 IleArgArgGlySerArgAspArgSerArgAlaSerArgSerArgSerAr 209
444 .....CATCTTCGTC 453
209 gglySerProLeuCysGlyAlaThrAlaThrSerCysProArgArgArg 226
454 AATGCGATGGACACCAATCCGCTGGCTGC..... 482
226 rgCysSerIleGlyAlaSerSerGlyCysProHisProValArgArg 242
483 CGACCCCTACGGTCATCATCAAGAAGCGCGGAGACTTCAACCGCGCC 532
243 SerProValAsnSerSerLysArgAlaHisArgCysThrAlaArgAr 259
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259 gglyArgPheArgGlyProThrSerArgAspThrGlyArgArgArgCysT 276
574 TGTAAAGCAGCAGCGGCGACGTCGCTGTAATAATGCTGCAATATCGA 623
276 rpArgTrpProArgProArgArgCys.....ArgCysSerArgArgTrp 290
624 AACACATGAATTTGGGCGCGCATCTCCGCGCTTGAGTGCACGCACA 673
291 GlyArgProLeuTrpAlaSerGlyCysProArgAlaArgTrp..... 304
674 TTCAATTTTCATCGACGTCGCGCGCAATAAAACCGTGTGGACCATCAAT 723
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724 TATCAAGACGTGATTGCTATCGACGCTTTGTCGTAACAGCGCTCTCAA 773
319 erProLysArgThrCysGlyArgArg...ValArgSerAspThrSerAla 334
774 TACCGACGCGGT..... 785
335 ArgArgSerArgCysProAlaSerSerProIleArgTrpThrGlyArgCy 351
786 .....GGTTGCTTGGGCGGCTGCAAGTCAAC 813
351 sArgArgTrpArgArgProLeuGlyCysSerProArg..... 363
814 AAACCGCGCTCTTGGTACCGTTGGTGGCAGAGGTGCTCAACTTAC 863
364 .....AlaThrCysThrAlaArgCysGlyArgAspGlyCysSerAlaPhe 378
863 ..... 863
379 pheGlyAsnProLeuHisArgSerLeuArgGlyProTrpAlaAlaProPh 395
864 .....CGCGGGAATTTGTTGACGCGGAC 888
395 eArgAlaHisArgSerArgSerThrThrArgCysAlaValArgGlyS 412

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162 laArgArgArgArg,..... 167
558 ACGTAAATCCATGCTGTAAAGCAGCGGCGAGCGTCCGCTCTGAAA 607
168 .....ArgArgArgArgGln..... 172
608 ATGCTGCCAATATCAACACATGAATTTGG.....CGGCCCGCAT 648
173 .....GlnPheArg.....**TrpArgGlyArgArgArgGlyA 184
649 CCTGCCG.....CTTGAGTGGCAGCGACATTCATTTCATCGA 686
184 rgCysArgArgAlaProLeuArgGlnTrpArgVal.....Arg 196
687 GCCAGTCGGCGCAATAAACCGGTGGACCATCAATTATCAAGACGTGA 736
197 ArgProArgArgSerArgHisGlyArgGlnHisArg..... 208
737 TTGCTATCGGAGCTTTGTTGCTAACAGCGCGCTGTGAATACCGAGCGGTG 786
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215 rgArgTrpGlyArgGlnAlaAspArg..... 223
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224 .....ProArgTrpArgArgArgCysArg.....ArgG 233
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250 GlyArgTrpArgProAlaileArg..... 257
1037 ACTCCATCAGCGGACCACTCTCGCCATTCCTAAACAACTCTTC 1086
257 ..... 257
1087 AAGTTCAGCAGCGCTCAACGGCGGCGACCGCCCATGTGTCGATCGG 1136
258 .....GlnArgArgArgArgProArgHis..... 266
1137 CACTTATAGCGCGTAATGCGGTGGACATCCTGCCTACCTT.....GC 1180
267 .....ArgArgAsnThrAlaGlyGlyGlyGluArgIleGlyAspG 280
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280 yPheValArgCysThrArgProThrArgArgHisArgProThrArgLeuA 297
1215 .....CGCGCAGGCTTTGGTGTGTAATTTGACGAGAGACCT 1256
297 laProIleAsnGluGlyPheGly...AlaGlyProGlyHisGlyHisPro 312
1257 .....CGCTTTGTGAGCTTCGCTCG.....CCGG 1282
313 LeuSerTrpArgSerGlyLeuArgTrpCysArgLysIleLeuProProG 329
1283 GCAATACGAATA 1295
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ID XX ABG05146 standard; Protein; 440 AA.
AC ABG05146;
DT 13-FEB-2002 (first entry)
XX Novel human diagnostic protein #5137.
DE Human: chromosome mapping; gene mapping; gene therapy; forensic;
KW food supplement; medical imaging; diagnostic; genetic disorder.
XX Homo sapiens.
OS WO200175067-A2.
PN 11-OCT-2001.
PD 30-MAR-2001; 2001WO-US08631.
PF 31-MAR-2000; 2000US-0540217.
PR 23-AUG-2000; 2000US-0649167.
XX (HYSE-) HYSEQ INC.
PA Drmanac RT, Liu C, Tang YT;
PI WPI; 2001-639362/73.
XX N-PSDB; AAS69333.
DR New isolated polynucleotide and encoded polypeptides, useful in
XX diagnostics, forensics, gene mapping, identification of mutations
PT responsible for genetic disorders or other traits and to assess
PT biodiversity -
XX Claim 20; SEQ ID No 35505; 103pp; English.
PS The invention relates to isolated polynucleotide (I) and
XX polypeptide (II) sequences. (I) is useful as hybridisation probes,
CC polymerase chain reaction (PCR) primers, oligomers, and for chromosome
CC and gene mapping, and in recombinant production of (II). The
CC polynucleotides are also used in diagnostics as expressed sequence tags
CC for identifying expressed genes. (I) is useful in gene therapy techniques
CC to restore normal activity of (II) or to treat disease states involving
CC (II). (II) is useful for generating antibodies against it, detecting or
CC quantitating a polypeptide in tissue, as molecular weight markers and as
CC a food supplement. (II) and its binding partners are useful in medical
CC imaging of sites expressing (II). (I) and (II) are useful for treating
CC disorders involving aberrant protein expression or biological activity.
CC The polypeptide and polynucleotide sequences have applications in
CC diagnostics, forensics, gene mapping, identification of mutations
CC responsible for genetic disorders or other traits to assess biodiversity
CC and to produce other types of data and products dependent on DNA and
CC amino acid sequences. ABG00010-ABG30377 represent novel human
CC diagnostic amino acid sequences of the invention.
CC Note: The sequence data for this patent did not appear in the printed
CC specification, but was obtained in electronic format directly from WIPO
CC at ftp.wipo.int/pub/published_pct_sequences.
XX Sequence 440 AA;
SQ
alignment_scores:
Quality: 125.00 Length: 480
Ratio: 0.654 Gaps: 26
Percent Similarity: 39.792 Percent Identity: 22.917
alignment_block:
US-09-303-518D-131 x ABG05146 ..
Align seg 1/1 to: ABG05146 from: 1 to: 440
56 TCATTATGACGGCGCGCATACCGATTCGCGTTCGTTGTCGCGAAGAA 105
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271	.SerTrpCysAlaArgTtrpAlaAlaThr***ProLeu.....ProS	284
921	TGCGATTGCAACAAGCGCGCATGATTATTATTTGGAGCGCTACCAACATCAGA	970
284	erAlaCysGlnArgLysTrpSerMetLeuTrpArgThrGlyThrAlaArg	300
971	TTTCCGTTATCGAAGAGCGCGCAGCAAGAGAGCTGTTCGGCTGGGTGGG	1020
301	ValSerThrSerAlaCysGlyAlaAlaProSerThrSerArgSer.....	315
1021	CCGCGACGGGACAAATACTCTCATCA...CGCGCACCACTCTCGGCCATT	1067
316	.ThrSerArgProSerThrProMetLeuArgAlaProValProAla...G	331
1068	CTTAATAAACAACCTCTTCAAGTTACACAGACGCCCTCAACGCGCGGAC	1117
331	lyTrpGlnProProAlaLeuHisProGlnProProArgProSerHisThr	347
1118	CGGCCATGGTACCGATCGGCACCTATGAGCGCGTAATCCGTGGACATC	1167
348	ArgGlnProTrpProSerAlaArgSer.....CysArgTrpArgTh	362
1168	CTGCCTACCTTGCTTTTCCGCGATTATTAATCTCGCGCATACCGACAGCG	1217
362	rCysThrThrArgProAlaSerSerThrSerSerProArgAlaThr***T	379
1218	GCAGGCTTTGG.....GTTGCTTGGATTGG.....	1243
379	hrSerHisTrpProProThrThrArgTrpArgMetSerArgCysSerThr	395
1244	...ACGAAGAAGACCTCGCTTTGTCAGCTTCGTCTGCCGGGCAAAATAC	1290
396	ProThrLysThrAsnCysThrCysMetArgGlyLeuGlyThrCysGlnAl	412
1291	GATACGGCCCGTGTTCGCAGAAAGTCTGCGAAACCA	1327
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seq_documentation_block:		
ID	AAV04955 standard; Protein; 573 AA.	
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AC	AAV04955;	
XX		
DT	06-JUL-1999 (first entry)	
XX		
DE	Mycobacterium species protein sequence 41T#3.	
XX		
KW	Secreted protein; Mycobacterium; primer; PCR; amplification; F	
KW	hybridisation; detection; vaccine; immunisation; infection.	
XX		
OS	Mycobacterium sp.	
XX		
PN	WO9909186-A2.	
XX		
PD	25-FEB-1999.	
XX		
PF	14-AUG-1998; 98WO-FR01813.	
XX		
PR	11-SEP-1997; 97FR-0011325.	
PR	14-AUG-1997; 97FR-0010404.	
XX		
PA	(INSP ) INST PASTEUR.	
PI	Gicquel B, Lim EM, Pellicic V, Portnoi D, Goguet de la salm	
PI	Guigueno A;	
DR	WPI; 1999-181045/15.	
DR	N-PSDB: AAX34206	

XX Mycobacterial DNA vectors containing reporter constructs - for  
PT identifying coding or promoter sequences involved in  
PT infection-associated protein expression  
XX  
XX Claim 32; Fig 41b; 309pp; French.  
XX  
XX Sequences AAY04742-Y05000 and AAY07201-Y07204 represent secreted  
CC proteins from various Mycobacterium species microorganisms. The  
CC encoding nucleotide sequences can be used as primers and probes for  
CC methods for detecting and identifying mycobacteria, especially belonging  
CC to the M. tuberculosis complex. The encoded proteins can be used in  
CC vaccines for immunisation against a bacterial or viral infection.  
XX  
SQ Sequence 573 AA:

alignment\_scores:  
Quality: 119.00 Length: 477  
Ratio: 0.626 Gaps: 20  
Percent Similarity: 39.832 Percent Identity: 23.270

alignment\_block:

US-09-303-518d-131 x AAY04955 ..

Align seg 1/1 to: AAY04955 from: 1 to: 573

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122 CCTCGATCAAAATCAAGAGGTGAACCGTCACAAAGG..... 161  
182 lArgAlaArgGlyValGlyArgCysGlyHisArgArgArg\*\*\*ArgGlyG 199  
162 .....CCAAAGTCTGTTGAAGACAAAGAAATCC 191  
199 lHisArgAlaGlyLysAspProArgThrAla\*\*\*ArgAlaArgArgCys 215  
192 GGGCGTAGTATTACTGGCGCGGCTTCAGCAAAATCGCGCTATTCCAC 241  
216 GlyArgGly.....GlyArgArgArgThrGlyPr 225  
242 GTGGCGAAAGCGGTACTTCAGTCAGTCGTGATTGCGGTTGAAGCAAC 291  
225 oAlaGlySerAlaGlyArgValAlaLeuHisHisLeuArgAlaGlyThrC 242  
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342 CAGCGAAAGTGGCGCGCAACCTGAT.....TC 370  
255 AspArgHisGlyTyrProThrProArgProAlaLeuArgGlyAspValSe 271  
371 AATCAGGCTTATGGACTCGCTTCGACCCGTCGCTTCAGCAAAATCCCT 420  
271 rValGlyGlyMet\*\*\*CysCysSerGlyGlyProVal...AlaGlySerT 287  
421 GCCGTAGATGCGGACCGCTTGCCTATCTTCGTCATGCGATGGACACCA 470  
287 hrGlnGlyIleGly\*\*\*ValGlyGlyHisArgArgCysSerAlaArgGln 303  
471 TCCGCTGGCTGCGGACCTACGGTCATCATCAAGAGCGCGCCGAGACT 520  
304 LeuLeuArgThrArgPro.....HisArgArgArgArgCysArgArgG 318  
521 TCAAACGGCGGCTGTGTATTGACCGCCCTGACCGAACGCTAAATCCAT 570  
318 ySerArgIleGlyGlyAlaSer\*\*\*ProAspArgAspLeuGly.... 333  
571 GTGTGTAAAGCAGCAGGCGCAGACGCTGCGGCTGAAATGCTGCCAATAT 620  
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334 .....AlaArgPhe 336  
621 CGAAACACATGAATT.....TGCGGGCCGCATCTCGCGCTTGA 661  
337 ArgAspGlnArgIleAlaGlyArgTrpLeuAspAlaGlyProArg..... 351  
662 GTGGCAGCACATTCATTTCATCGAGCGAGTCGGCGGGAATAAAACCGTG 711  
352 .....ArgAlaGlyGlyArg..... 357  
712 TGGACCATCAATATCAAGACGTGATTCTATCGAGCTTTGTTTCGTAAC 761  
358 .....ArgArgArgCysArgArgAla..... 364  
762 AGGCGCTCTGATACCGAGCGGTGCTTGGCGGCGCTGCAAGTCA 811  
365 .....ValArgArgGlyArgLeuArgAlaAlaThrGlySe 377  
812 ACAACCGCGCTCTTGTGCTACCGCTTTGGGTGC.....GAAG 849  
377 rArgArgArgAspThrGlyArgArgTyrGlnCysProProAlaGlyAlaG 394  
850 GTGTCTCAACATTACCGCGCGCAATTGTTGACGCGGACACCGCGTGAT 899  
394 lArgGlyArgHisArgArgAla..... 402  
900 TTCCGCTTCGCTATTGAACGCTGCGATTGCACAGCGCGCATGATTATT 949  
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417 yGlyArgValTyrArgGlyAsp...ArgLeuGlyArgArgArgGlyThrA 433  
1000 GAGCTTTCGCTGGGTTCGCGCGCAGCGGCAAAATA..... 1037  
433 rGAlaAspArgIleAspGlyAlaGlyValGlyArgAlaGlyArgAla\*\*\* 449  
1038 .....CTCCATCAGCGCGCACCATCTCGC 1060  
450 ArgGlyProProGlyArgArgArgLeuGlnHisGlyProCysArgArg 466  
1061 GCCATTTCTATAAAACAACCTCTCAAGTTCCAGCAGC..... 1100  
466 gCysPheProAlaArgIleGlyAlaHisCysHisProLysGlyAlaAspL 483  
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500 GlyAlaAspArg..... 503  
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504 .....ArgArgArgCysArgArgGlyGlyHisA 513  
1216 GCGCAGCGCTTGGGTGCTTGGAAATGGAGAGAACCTCGCTTTGTG 1265  
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1266 CAGCTTCGT.....CTCCCGCGGCAATACG 1291  
530 AsnTrpArgArgArgAsnArgArgArgGlyCysArgProGlyThrAlaCy 546  
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546 sAlaArgProProSerArgHisArgAlaGly 556

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seq\_documentation\_block:



XX DE Novel human diagnostic protein #3722.  
XX KW Human; chromosome mapping; gene mapping; gene therapy; forensic;  
XX KW food supplement; medical imaging; diagnostic; genetic disorder.  
XX OS Homo sapiens.  
XX PN WO200175067-A2.  
XX PD 11-OCT-2001.  
XX PF 30-MAR-2001; 2001WO-US08631.  
XX PR 31-MAR-2000; 2000US-0540217.  
XX PR 23-AUG-2000; 2000US-0649167.  
XX PA (HYSE-) HYSEQ INC.  
XX PI Drmanac RT, Liu C, Tang YT;  
XX DR WPI; 2001-639362/73.  
XX DR N-PSDB; AAS67918.  
XX CC New isolated polynucleotide and encoded polypeptides, useful in  
PT diagnostics, forensics, gene mapping, identification of mutations  
PT responsible for genetic disorders or other traits and to assess  
PT biodiversity -  
XX PS Claim 20; SEQ ID No 34090; 103pp; English.  
XX CC The invention relates to isolated polynucleotide (I) and  
CC polypeptide (II) sequences. (I) is useful as hybridisation probes,  
CC polymerase chain reaction (PCR) primers, oligomers, and for chromosome  
CC and gene mapping, and in recombinant production of (II). The  
CC polynucleotides are also used in diagnostics as expressed sequence tags  
CC for identifying expressed genes. (I) is useful in gene therapy techniques  
CC to restore normal activity of (II) or to treat disease states involving  
CC (II). (II) is useful for generating antibodies against it, detecting or  
CC quantitating a polypeptide in tissue, as molecular weight markers and as  
CC a food supplement. (II) and its binding partners are useful in medical  
CC imaging of sites expressing (II). (I) and (II) are useful for treating  
CC disorders involving aberrant protein expression or biological activity.  
CC The polypeptide and polynucleotide sequences have applications in  
CC diagnostics, forensics, gene mapping, identification of mutations  
CC responsible for genetic disorders or other traits to assess biodiversity  
CC and to produce other types of data and products dependent on DNA and  
CC amino acid sequences. ABG00010-ABG30377 represent novel human  
CC diagnostic amino acid sequences of the invention.  
CC Note: The sequence data for this patent did not appear in the printed  
CC specification, but was obtained in electronic format directly from WIPO  
CC at ftp.wipo.int/pub/published\_pct\_sequences.  
XX SQ Sequence 696 AA;

alignment\_scores:  
Quality: 115.50 Length: 462  
Ratio: 0.553 Gaps: 23  
Percent Similarity: 45.238 Percent Identity: 24.459  
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77 TTACCGAAGTCGGTGTGTCGCGNAGATATGTCGGCATCGGCCCTC. 125  
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25 sasparg\*\*\*Arg.....ArgValSerProThrA 35

126 .....GATCAAAATCAAGGAAGTGAAGCGCTCAA 155  
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35 rgSerGlyLysArgArgGlyAlaGluGluLysAsnArgGlnGluLysLys 51  
156 AAAGGCCCAAGTGTCTTTTGAAGACAAAGAAATCCGGGCGGTAGTATTTA 205  
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52 LysGlyArgGluLysGluArgGluLysArgSerGluArgGln..... 66  
206 CTGCGCCCGCTTCAGGCAAAATCCCGCTATTACCGCTGGCGGAAAGCGC 255  
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67 .....ArgaspArgArgArgGlySgluGlnArgLys..... 77  
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78 .....GluGluGlnArgArgAlaArgThrAsnGluArg 89  
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90 LysProArgGlnThrGlnAlaAsnGlyAlaThrSerSer\*\*\*LysAlaSe 406  
356 GCGCGCAACCTGATTCAATCAGGCTTATGAGCTGCGCTTCGCACCGCTCCG 405  
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106 rAlaGlnGlnAlaGlyMet.....TrpGlyLysSerPro\*\*\*T 419  
406 TTACGCAAAATCCCTGCGCTAGATCCGAGCGCTTCGCCATCTTCGTC 455  
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119 hrAspAlaThrAlaIleArgArgGlyGlyAlaProCysSerSerArgArg 455  
456 TCGGATGGACACCAATCCGCTGGCTGCGGACCGCTACGCTCATCATCAA 505  
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136 ThrCysLeuAsnGlnGlyThrIleAlaThrProSerGlyArg..... 505  
506 AAGCGCGCAAGACTCAAAACGCGCTGTTGGTATTGAGCGCGCTGACC 555  
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150 ....ArgArgHisGlyAspAlaGly\*\*\*ProGlyLeuAlaSerGluHis 555  
556 GAACGTAAATCCATGTGTGTAAAGCAGGCGGCGGAGCGCTGCTCA 605  
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165 AspAlaSerGlyHisGlyCysLeuArgThrGlyAlaGly\*\*\*ProSerAs 605  
606 AAAT.....GCTGCCAATATCGAAACAC 628  
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181 psThrGluSerValCysArgArgProLeuAlaMetHisValProThrH 198  
629 ATGAATTTGCGCGCGCATCTCGCGCTGAGTGGCAGCGACCATTCAT 678  
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198 isGluSerHisGlyPro.ValPheThrArgLeuValSerHisThrPheHi 678  
679 TTCATCGCGCGAGTCGCGCGCAATA...AAACCG.....TGTGACCAT 719  
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897 alCysThrThrIleValGluThrGlyLeuAspIleSerAsnAlaAsnThr 913
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1191 ArgGly 1192
seq_name: /SIDS1/gcgdata/geneseq/geneseq-emb1/AA2001.DAT: AAB96063
seq_documentation_block:
ID AAB96063 standard; Protein; 822 AA.
XX AAB96063;
XX AC
XX XX
DT 29-OCT-2001 (first entry)
XX XX
DE Putative P. abyssi pyruvate kinase #1.
XX XX
KW Hyperthermophilic archaeon; hyperthermophilic protein.
XX OS
XX PY Pyrococcus abyssi.
XX FR FR2792651-A1.
XX PD 27-OCT-2000.
XX PF 21-APR-1999; 99FR-0005034.
XX PR 21-APR-1999; 99FR-0005034.
XX XX
XX (CNRS ) CNRS CENT NAT RECH SCI.
XX (IFRE-) IFREMER INST FR RECH EXPL MER.
XX PI
XX PI Forterre P, Thierry JC, Prieur D, Dietrich J, Lecompte O;
XX PI Querrelou J, Weissenbach J, Saurin W, Heilig R;
XX WPI; 2001-126236/14.
XX New nucleotide sequences isolated from Pyrococcus abyssi encode
XX proteins useful in industry -
XX Claim 7; Pages 677-680; 1657pp; French.
XX The present invention relates to the genomic sequence of Pyrococcus
XX abyssi (see AAF86431 and AAH41223-7) and P. abyssi proteins. P. abyssi is
XX a hyperthermophilic archaeon, which is isolated from deep-sea
XX hydrothermal vents. The present sequence is one such P. abyssi protein.
XX The proteins of the present invention have various potential industrial
XX uses, since the proteins are stable at very high temperatures, some up to
XX 110 degrees centigrade.
XX Note: This patent is in the same patent family as WO2000065062, which
XX contains additional sequences as shown in AAB99132-AAB99143,
XX AAH75903-AAH75920 and AAG66436.
XX Sequence 822 AA;
SQ
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alignment\_scores:
Quality: 113.00 Length: 475
Ratio: 0.467 Gaps: 20
Percent Similarity: 50.947 Percent Identity: 20.421
alignment\_block:

US-09-303-518D-131 x AAB96063

Align seg 1/1 to: AAB96063 from: 1 to: 822

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31 CCATCCGCGGAGACCGGAGCAAGTCAATTATGAC..... 66
|||||:|||||:|||||:|||||:|||||:|||||:|||||:
238 ProValThrAsnAsnArgAsnGluIleMetIleAsnAlaSerTrpGlyLe 254
67 .GCCCGCGCCATTACCGAAGTCCGTTGCGGAGGAATATGTCGGCA 115
|||||:|||||:|||||:|||||:|||||:|||||:|||||:
254 uGlyGluAlaValValSerGlyAlaValThrProAspGluTrpIleValG 271
116 TGGCCCGCCCGTGAATAAAGAGGAGTGAAGCCGTCAAAAGAGCCAA 165
|||||:|||||:|||||:|||||:|||||:|||||:|||||:
271 luLysGlyThrTrpLysIleLysGlu...LysValIleAlaLysLysGlu 286
166 GTCTCTTTTGAACACAAAGATCCGCGGTAGTATTACTGCGCGCGC 215
|||||:|||||:|||||:|||||:|||||:|||||:|||||:
287 ValMet.....ValIleArgAsnProG1 294
216 TTCAGGCAAA.....ATCGCGCTATTACCGTGGC... 246
|||||:|||||:|||||:|||||:|||||:|||||:|||||:
294 uThrGlyLysGlyThrValThrValLysValAlaGluTrpLeuGlyProG 311
247 .....GAAGCCGCTACTT.....CAGTCAGTCGTGATGCC 279
|||||:|||||:|||||:|||||:|||||:|||||:|||||:
311 luTrpValGluLysGlnValLeuThrAspGluGlnIleIleGluValAla 327
280 GTTGAAGCAACGACCAAAATCAGTTCGACGCTACGTACCTGAAGCGCT 329
|||||:|||||:|||||:|||||:|||||:|||||:|||||:
328 LysMetGlyGlnLysIleGluGluHisTrpGlyTrpProGlnAspIleG1 344
330 GCGAAATGACAGCGGAAAAAGTGGCGGCAACCTGATTCAATCAGGCT 379
|||||:|||||:|||||:|||||:|||||:|||||:|||||:
344 uTrpAlaTrpAspLysAspGlyLysLeuTrpIleValGlnSerArgp 361
380 TATGAGTCGCTGTCGACCCGCTCGCTCAGCAAAATCCCTCGCTAGAT 429
|||||:|||||:|||||:|||||:|||||:|||||:|||||:
361 rovalThrThrLeuLysGluThrThrThrGluGluVal... 376
430 GCCGAGCGCTGCTCCATCTTCATCGATGCGAGACCAATCCG...CT 476
|||||:|||||:|||||:|||||:|||||:|||||:|||||:
377 ...GluGluAlaGluValIleLeuLysGlyLeuGlyAlaSerProGlyI1 392
477 GCCTCGGACCTACGCTCATCATCAAGAGCCGCGCAAGACTTCAAAC 526
|||||:|||||:|||||:|||||:|||||:|||||:|||||:
392 eGlyAlaGlyArgValValIlePheAspAlaSerGlu..... 405
527 GCGGCTGTGTGATTGAGCGCCCTGACCGACGTAAATCCATGCTGTGT 576
|||||:|||||:|||||:|||||:|||||:|||||:|||||:
406 .....IleAspLysValLysGlyGluGlyAspValLeuValThr 417
577 AAAGCAGCAGCGGACAGAC...GTGCGCT...GAAATGCTGCAATAT 620
|||||:|||||:|||||:|||||:|||||:|||||:|||||:
418 ThrMetThrAsnProAspMetValProAlaMetLysArgAlaAlaI1 434
621 CGAACACATGAATTTGCGCGCCGCTCCTCGCGGCTTGTAGTGGCAGC 670
|||||:|||||:|||||:|||||:|||||:|||||:|||||:
434 elleThrAspGluGlyGlyArgThrSerHisAlaAlaIleValSerArg 451
671 ACATTTCATTCATCAGCAGCGCGGGAATAAACCGCTGAGCCATC 720
|||||:|||||:|||||:|||||:|||||:|||||:|||||:
451 luLeuGlyIleProAlaValValGlyThrLysGluAlaThrLysLysLeu 467
721 AATTATCAAGACGATGTCATCGGA..... 747
|||||:|||||:|||||:|||||:|||||:|||||:|||||:
468 LysThrGlyAspTrpValThrValAspGlyThrArgGlyLeuValTyrLy 484
748 .....CGTTGTTGCTAAACAGCCCTCTCAATACCGAGCGCGTGG 787
|||||:|||||:|||||:|||||:|||||:|||||:|||||:
484 sGlyIleValLysSerLeuValGluLysLysLysLysLysGluAlaAla 501
788 TTGCTTTGGCGGCGCTGCAAGTCAACAAACGCGCCTCTTTCGTACCGTT 837
```

```
|||||:|||||:|||||:|||||:|||||:|||||:|||||:
501 laAlaProGlyAlaAlaValAlaAlaProLeuValThrGlyThrLeu 517
838 TTGGTGTGGAAGGTGCTCAACTTACCGCGCGCAATTTGGTTGACGGGA 887
|||||:|||||:|||||:|||||:|||||:|||||:|||||:
518 ValLysValAsnValSerMetProGluValAlaGlu..... 529
888 CAACCGCTGATTTCCGGTTCGGTATTGAACGGTGGATTGCACAAGCGC 937
|||||:|||||:|||||:|||||:|||||:|||||:|||||:
530 ...ArgAlaAlaThrGlyAlaAspGlyValGlyLeuLeuArgAlaG 545
938 CGCATGATTATTGGACGCTACCAATCATCATTCCTGTTATCGAAGAA 987
|||||:|||||:|||||:|||||:|||||:|||||:|||||:
545 luHisMetIleLeuSerIleGlyGlnHisProValLysPheIleLysGlu 561
988 GCGCGCACAAAGAGCTGTTGCGGTGCGCGCGACGCGGACAAATA 1037
|||||:|||||:|||||:|||||:|||||:|||||:|||||:
562 GlyLysGluGluLeuValGluLysLeuAlaGluGlyIleGluLysVa 578
1038 CTCATACGCGCGCACCACTCTCGGCCATTT..... 1067
|||||:|||||:|||||:|||||:|||||:|||||:|||||:
578 laAlaAla..AlaPheTyrProArgProValTrpTyrArgThrLeuAspAl 594
1068 .CCTAAAAACAACTCTTCAAGTT.....CACGACAGCGCTCAAC 1107
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594 aProThrAsnGluPheArgGluMetProGlyGlyGluAspGluProGluG 611
1108 GCGCGGACCGCGCATGGTACCGATCGGCACCTTATGAGCGCGTAATGCC 1157
|||||:|||||:|||||:|||||:|||||:|||||:|||||:
611 luArgAsnProMetLeuGlyTrpArgGly.....IleArgArgGly 624
1158 GTTGGACATCTCCTACCTTGTGTTGCGGATTTAATCTCGCGCGATA 1207
|||||:|||||:|||||:|||||:|||||:|||||:|||||:
625 LeuAspGlnProGluLeuLeuArg...AlaGluPheLysAlaIleLysLy 640
1208 CCGACAGCGCGCAGGCTTGGTTGCTTGGAAATGGACGAAGAAGACCTC 1257
|||||:|||||:|||||:|||||:|||||:|||||:|||||:
640 sValValGluLysGlyTyrAsnAsnIleGly..... 650
1258 GCTTTGTCAGCTTCGTCG.....CCGCGGCAATACGAATACGG 1298
|||||:|||||:|||||:|||||:|||||:|||||:|||||:
651 .....ValMetLeuProLeuValSerHisProGluGlnIleArgLysAla 665
1299 CCCGCTGTTCGCAAGTGTGGA 1322
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666 LysGluIleAlaArgSerValGly 673
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seq\_name: /SIDS1/gcgdata/geneseq/geneseq-emb1/AA2001.DAT:AAB59817

seq\_documentation\_block:

ID AAB59817 standard; Protein; 999 AA.

XX AAB59817;

XX AC AAB59817;

XX DT 04-APR-2001 (first entry)

XX DE TutD protein #8.

XX KW Toluene degradation; enzyme; waste degradation; TutD.

XX OS Thauera aromatica.

XX OS Xanthomonas maltophilia.

XX OS Geobacter metallireducens.

XX OS Azorarcus toluolyticus.

XX PN WO200072650-A2.

XX PD 07-DEC-2000.

XX PF 24-MAY-2000; 2000WO-US14298.

XX PR 01-JUN-1999; 99US-0323872.

XX

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alignment_scores:
  Quality: 112.50      Length: 536
  Ratio: 0.516        Gaps: 26
  Percent Similarity: 40.672  Percent Identity: 23.694

alignment_block:
US-09-303-518D-131/rev x AAB59817  ..

Align seg 1/1 to: AAB59817 from: 1 to: 999

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112 CysThrSerArgGlyArgSerArgCysSerProAspArgCysCysArgTr 128
1273 .CGAAGCTGCAACAGCGAGGTCTCTTCGTGCCAATTCACAGCAACCCA..1226
128 pSerArgCysSerSerProSerProArgArgCysProProSerSerProA 145
1225 .....AAGCCTGCGGGCTGCGGTATCGCGACGATTAAATCGCGCAAAA 1181
145 laGlyAlaProGlyAlaThrCysSerArgArgProPheSerArg..... 159
1180 GCAAGTAGGAGAGATGTCACAGCGCATTCACGCTCATTAAGTCGCGATC 1131
160 SerArgaspSerAlaGlyProArgAlaAlaArgPheArgArgCysArgAs 176
1130 GGTACCATGGCGGCTGCGCGCGGTGTGACGGCTGTCGTGAACCTGAAGAG 1081
176 paLacCysGluArgArgAlaArg..... 183
1080 TTGTGTTTTTAGGAAATGCGCGAGAGTGGTGGCGGTGATGGAGTATTGT 1031
184 .....CysProGlyProArgSerAla 190
1030 CCGGTGCGGGCGCAACCCAGCGCAACAGACTTTTGTGCGCGGCTTCTTCG 981
191 ProSerIleArgArgGlySerArgaspArgSerArgAlaSerArgSerAr 207
980 ATAACGGAAATCTGAT...TGTTGTAGCTGCCAAATATCATCGCGGCC 934
207 gSerArgGlySerProLeuCysGlyAlaThrAlaThr..... 219
933 TTGTGCAATCGCACCGTTCATACCGCAACCGGAAA..... 899
220 .....SerCysProArgArgArgArgCysSerIleGly 230
988 ...TCACGGGTTGTCGCGGTCAACCAATTCGCGCGGTAACTTGAGAC 852

```

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157 .....TTTTACGGCTTCACCTTCCTTGATTTT 130
158 oSerSerProLeuAsnArgProPheAlaArgSerAlaPro.....A 531
129 CATCGAGGCGGATCGGCACATATCTTCGCCAAGCAAGCGGACTCGG 80
130 :||||:||||:||||:||||:||||:||||:||||:||||:||||:
531 laSerThrProCysArgArgHisAsnArgArgTyrGlySerArgArg 547
79 .....TAATGGCGGGCGCGTCATAAA..... 59
548 ProPheArgArgPheAlaCysSerTyrSerSerGlnHisAspProAl 564
58 .....TGACTTGCTCGGCTCGCCCGC 37
564 aSerGlnAspProGlnArgGlyThrCysProLeuArgAsnAlaCysProG 581
36 GATGGGCA 29
581 lyTrpAla 583
seq_name: /SID1/gcgdata/geneseq/geneseq-emb1/AA1999.DAT.AAY28654
seq_documentation_block:
ID AAY28654 standard; Protein: 802 AA.
XX AC AAY28654;
XX DT
XX DE
XX DW
XX KW Murine Cytoplasmic phosphatase, Z70PEP protein.
XX KW Lymphoid protein Tyrosine Phosphatase; Lyp protein; lymphoid cell;
XX KW intracellular tyrosine phosphatase; ptpase; lymphocyte; murine;
XX KW protein tyrosine kinase; PTKs; immunosuppressant; PEST sequence;
XX KW T cell antigen receptor signalling; autoimmune disease; transplant;
XX KW cytokine receptor signalling.
XX OS Mus sp.
XX FH
XX Key Location/Qualifiers
XX Domain 27..288 /label= ptpase_domain
XX FT /note= "Catalytic protein tyrosine phosphatase domain"
XX FT Binding-site 613..621 /label= SH3_binding_site
XX FT /note= "SH3 binding site"
XX FT Binding-site 689..695 /label= SH3_binding_site
XX FT /note= "SH3 binding site"
XX FT Binding-site 790..798 /label= SH3_binding_site
XX FT /note= "SH3 binding site"
XX PN WO936548-A1.
XX PD
XX XX
XX XX 22-JUL-1999.
XX PF 18-JAN-1999; 99WO-CA00038.
XX XX
XX PR 16-JAN-1998; 98CA-2220853.
XX XX (HSC-) HSC RES & DEV LP.
XX PA
XX XX
XX PI Roifman CM;
XX XX
XX XX WPI; 1999-444404/37.
XX XX
XX PT New nucleic acid encoding intracellular tyrosine phosphatase and
XX PT related proteins, used to modulate signalling through T cells,
XX PT particularly as immunosuppressant
XX PS
XX PS Disclosure; Page 63-64; 105pp; English.

```

```

XX The present protein sequence is that of the murine phosphatase, Z70PEP
CC that has a single catalytic domain. The non-catalytic portion of the
CC phosphatase contains unique sequences, including five PEST sequences
CC rich in Pro, Glu or Asp, Ser and Thr. Z70PEP shares about 70% sequence
CC identity with the human cytoplasmic phosphatase LypL. Lyp proteins are
CC important for regulation of T cell antigen and cytokine receptor
CC signalling and for early and late stages of T cell differentiation.
CC Z70PEP has immunosuppressive activity. Compounds that increase
CC expression of Lyp protein can be used as immunosuppressive agents to
CC reduce or prevent T cell activation or proliferation, to control
CC thymocyte differentiation, to treat autoimmune diseases and transplant
CC situations.
XX
XX Sequence 802 AA;

```

```

alignment_scores:
Quality: 112.00 Length: 351
Ratio: 0.622 Gaps: 20
Percent Similarity: 51.282 Percent Identity: 24.501
alignment_block:
US-09-303-518D-131/rev x AAY28654 ..
Align seg 1/1 to: AAY28654 from: 1 to: 802

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955 GTCCCAATATCAT...GCGCGCTTGTGCATTCGCACCGTTCATATACC 909
956 :|||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:
957 IleSerAspAsnHisLeuGlyArgGluIleGlnAlaGlnCysSerIlePr 313
958 GAACCGG.....AAATCACGCGGTGTTCGCGCTCAACCAATTC 871
959 :||||:||||:||||:||||:||||:||||:||||:||||:
960 oGluGlnSerLeuThrValGluAlaAspSerCysProLeuAspLeuProL 330
961 :||||:||||:||||:||||:||||:||||:||||:||||:
962 GCGCG.....CGTAAAGTTGACACACCTTCG...CACCCAAACGCGTAC 830
963 :|||:||||:||||:||||:||||:||||:||||:||||:
964 ysAsnAlaMetArgAspValLysThrThrAsnGlnHisSerLysGlnGly 346
965 :|||:||||:||||:||||:||||:||||:||||:||||:
966 GCAAGAGCGCGGTGTTCGCTGCTGACGCGCGCCGCAAGCAACCGCGCG 780
967 :|||:||||:||||:||||:||||:||||:||||:||||:
968 Ala.GluAlaGlu.....SerThrGlyGlySerSerL 357
969 TCGTATTACAGCGGCTGTTCAGCAACAGCTCCGATAGCAATACGTC 730
970 :||||:||||:||||:||||:||||:||||:||||:||||:
971 euGlyLeuArgThrSer.....ThrMetAsnAlaGluGluGlu 369
972 :||||:||||:||||:||||:||||:||||:||||:||||:
973 TTGATAATTGATGTCACACGCGTTTATTCGCGCGCGCTGCTCGATGA 680
974 :||||:||||:||||:||||:||||:||||:||||:||||:
975 LeuValLeuHisSerAlaLysSerSerProSerPheAsnCysLeu...G 385
976 :||||:||||:||||:||||:||||:||||:||||:||||:
977 AATCAATGTGCGTGC...ACTCAAGCGCGCGGCA 648
978 :||||:||||:||||:||||:||||:||||:||||:||||:
979 uLeuAsnCysGlyCysAsnAsnLysAlaValIleThrArgAsnGlyGlnA 402
980 :||||:||||:||||:||||:||||:||||:||||:||||:
981 TCGCGCGCGCGC.....AAATTCATGTGTTTC 622
982 :||||:||||:||||:||||:||||:||||:||||:||||:
983 laArgAlaSerProValValGlyGluProLeuGlnLysTyrGlnSerLeu 418
984 :||||:||||:||||:||||:||||:||||:||||:||||:
985 GATATTGGCAGCATTTTCAGACGCGCACGCTGCTGCGCTGCTGTTCACA 572
986 :||||:||||:||||:||||:||||:||||:||||:||||:
987 AspPheGlySerMetLeu.PheGly...SerCysProSerAlaLeuPro. 433
988 :||||:||||:||||:||||:||||:||||:||||:||||:
989 CATGGATTTTACGTCAGCGGCTCAATACCAACAGCGCG..... 528
990 :|||:||||:||||:||||:||||:||||:||||:||||:
991 .....IleAsnThrAlaAspArgTyrHisAsnSerLysGlyProValLys 448
992 :||||:||||:||||:||||:||||:||||:||||:||||:
993 CGTTGAAGTCTTCGCGGCTCTTTTGATG..... 498
994 :||||:||||:||||:||||:||||:||||:||||:||||:
995 ArgThrLysSerThrProPheGluLeuIleGlnArgLysThrAsnAs 465
996 :||||:||||:||||:||||:||||:||||:||||:||||:
997 .ATGACCGTATGGTCGCGCAGCGGATGTTGTGTC..... 462
998 :||||:||||:||||:||||:||||:||||:||||:||||:

```



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1009 CGAACAGCTCTTTGCTGCGCGCTTCTTCGATACACGGAAATCTGATTGTGG 960
||||| :|
210 roAsn...LysCysTyrAlaProSerAlaIlePro..... 220
959 TAGCTGCCAATAATATCATCGCGCTTGTGGAATCGCACCGCTTCAATAC 910
||||| :|
221 ...ThrProGlnArgThrSerThrPro...GlyLeuAlaLeuPheProG1 235
909 CGAACCGGAAATCACCGCGTGTTCGCGCNCAC 876
||||| :|
235 yProProSerProValAlaAsnSerThrProLeuThrLeuProVal 252
875 ..AATTGCGCG .....GCGTAAGTTGAGACACCTTCGCACCCAAA 837
||||| :|
252 alGlnSerProLeuAlaThrAlaAlaSerAlaSerThrSerAlaProVal 268
836 ACGGTACGCAAGAGCGCGTGTGTGACTTCGACGCGCCGCCAACGCAAC 787
||||| :|
269 SerCysGlySerSerAlaSerLeuLeuArgGlyProHisProGlyThrSe 285
786 CACGCGCTCGGTATTACAGCGCGCTTACGACAAACGTCGATAGCAA 737
||||| :|
285 rAspLeuHisIleSerSerThrProAlaAlaThrThrLeuProVal.... 300
736 TCACGTCTGTGATTAATGATGTCACAGCGTTTATTGCGCGCGACTGGC 687
||||| :|
301 .....MetIleLysThrGluProThrSerProThrPro 311
686 TCGATGAATGAATGTCGCTGCCACTCAAG...CGGCGAGGATGGGGCC 640
||||| :|
312 Ser.....AlaPheLysGlyProSerHisSerGlyAs 322
639 GCCAATTCATGTGTTTCGATATGCGACGATTTTCAGACGCGAGCTG 590
||||| :|
322 nProSerHisGlyThrLeuGlyLeuSer.....GlyThrLeuG 335
589 CGCTGCTGCTTTACACACATGGAATTTACGTTGCGTCAGCGCTCAAT 540
||||| :|
335 lYArgAlaTyrThrSerThr..... 341
539 ACCAACAGCGCGCTTGAAGTCTTCGCGCGCTTCTTGTGATGATGACG 490
||||| :|
342 .....SerValProIleSerLeuSerAlaCysLe 351
489 AGGTGCGCACCGCGGATTTGTCATCGCATTCAGCAAGATGGCGA 440
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351 uAsnProAlaLeuSerGlyLeuSerSerSerThrPro.....LeuA 366
439 ACGCTCGGCATCTACGCGAGGATTTGCTGAACGCGGCGGTGCGAAGC 390
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366 sNGLySerAsnProLeuSerSerIleSerLeuProProHisGlySerSer 382
389 CAGTCCATAGCCTGATTGAATCAGTTGCGGCGCCTTTTCGCTGCT 340
||||| :|
383 ThrProIleAlaProValPheThrAlaLeuProSerPheThrSerLeuTh 399
339 CAATTTGCCAGCGCTTACGTAGCTAGCGTTCGAACCTCGATTTCGTCGT 290
||||| :|
399 rAsnAsnPheProLeuThrGlyAsnProSerLeuAsnProSerValSerL 416
289 TGCCTTCAACGCAATCAGCACTACGTAAGTACGCGCTTTTCG..... 246
||||| :|
416 euProGlySerLeuIleAlaThrSerSerThrAlaAlaThrSerThrSer 432
245 .....CCACGGTGAATAGCGCGATTTTG..... 222
433 LeuProHisProSerThrAlaAlaValLeuSerGlyLeuSerAlaSe 449
221 ....CTGAACCGCGCA..... 207
449 rAlaProValSerAlaAlaProPheProLeuAsnLeuSerThrAlaValP 466
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seq\_documentation\_block:

ID AAY31745 standard; Protein; 430 AA.

XX AAY31745;

XX 22-NOV-1999 (first entry)

XX Mycobacterium tuberculosis specific DNA-encoded polypeptide.

XX Tuberculosis; infection; diagnosis; DNA probe.

XX Mycobacterium tuberculosis.

XX Key Location/Qualifiers

FT Misc-difference 4 /note= "encoded by TGA"

FT Misc-difference 6 /note= "encoded by TGA"

FT Misc-difference 20 /note= "encoded by TGA"

FT Misc-difference 29 /note= "encoded by TGA"

FT Misc-difference 54 /note= "encoded by TGA"

FT Misc-difference 64 /note= "encoded by TGA"

FT Misc-difference 69 /note= "encoded by TGA"

FT Misc-difference 89 /note= "encoded by TGA"

FT Misc-difference 99 /note= "encoded by TGA"

FT Misc-difference 114 /note= "encoded by TGA"

FT Misc-difference 119 /note= "encoded by TGA"

FT Misc-difference 129 /note= "encoded by TGA"

FT Misc-difference 159 /note= "encoded by TGA"

FT Misc-difference 169 /note= "encoded by TGA"

FT Misc-difference 182 /note= "encoded by TAG"

FT Misc-difference 185 /note= "encoded by TGA"

FT Misc-difference 219 /note= "encoded by TGA"

FT Misc-difference 259 /note= "encoded by TGA"

FT Misc-difference 269 /note= "encoded by TGA"

FT Misc-difference 291 /note= "encoded by TGA"

FT Misc-difference 323 /note= "encoded by TGA"





```

297 gCysArgGluCysCysAsnCysArgCysTrpArgCysArgGluCysSera 314
934 GCGCGCATGATTATTGGG.....ACG 956
||||| |||
314 rArgProGlyLeuProGlyArgAsp***ArgProValGlyHisArgLys 330
957 CTACCAACATCAGATTTCGTTTATCGAAGAGCGCGCAGCAAGAGCTGT 1006
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331 IleProThrCysCysPheArgCys***ArgSerProArgSerArgProAl 347
1007 TCGGCTG.....GGTTCGCCCGCAGCGCGGACAAATCTCCATCAG 1047
:|Leu***TrpProGlySerCys***ThrAsnProIleArgCysCysP 364
1048 CGCACCACCTCGCGCATTCCTTAAACAACTCTTCAAGTTCACGAC 1097
364 roSer***SerArgProIleProAlaArgProArgLeu.....Pro 377
1098 AGCGGTACAGCGCGGCGACCG.....CGCCATGTACCGATCGGCACTT 1141
:||||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
378 GlyArgSerTyrArgTrpProThrLysSerGlyArgSerGlnAsnCy 394
1142 ATGAGCGCTAATGCGTTGGACATCTCCCTACCTTGTCTTTGCGCGAT 1191
394 strPhisArgSer.....G 400
1192 TTAATCGTCGGGATACCGACGCGC 1217
400 lySerArgThrArgTyrArgHisArg 408

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seq\_name: /SIDS1/gcgdata/geneseq/geneseq-emb1/AA1999.DAT:AAV04954

seq\_documentation\_block:

ID AAV04954 standard; Protein: 572 AA.

XX AC AAV04954;

XX DT 06-JUL-1999 (first entry)

XX DE Mycobacterium species protein sequence 41T#2.

XX KW Secreted protein; Mycobacterium; primer; PCR; amplification; probe;  
XX KW hybridisation; detection; vaccine; immunisation; infection.  
XX OS Mycobacterium sp.  
XX FN W0909186-A2.  
XX PD 25-FEB-1999.  
XX PF 14-AUG-1998; 98WO-FR01813.  
XX PR 11-SEP-1997; 97FR-0011325.  
XX PR 14-AUG-1997; 97FR-0010404.  
XX (INSP ) INST PASTEUR.  
XX GI Gicquel B, Lim EM, Pellicic V, Portnoi D, Goguet de la Salmoniere Y;  
XX PI Guigueno A;  
XX WPI; 1999-181045/15.  
XX DR N-PSDB; AAX34206.  
XX PT Mycobacterial DNA vectors containing reporter constructs - for  
XX PT identifying coding or promoter sequences involved in  
XX PT infection-associated protein expression  
XX PS Claim 32; Fig 41T; 309pp; French.  
XX Sequences AAV04742-Y05000 and AAV07201-Y07204 represent secreted  
XX CC proteins from various Mycobacterium species microorganisms. The  
XX CC encoding nucleotide sequences can be used as primers and probes for  
XX CC methods for detecting and identifying mycobacteria, especially belonging

CC to the M. tuberculosis complex. The encoded proteins can be used in  
CC vaccines for immunisation against a bacterial or viral infection.

XX SQ Sequence 572 AA;

alignment\_scores:  
Quality: 108.50 Length: 507  
Ratio: 0.580 Gaps: 31  
Percent Similarity: 36.884 Percent Identity: 21.105

alignment\_block:

US-09-303-518D-131 x AAV04954 ..

Align seg 1/1 to: AAV04954 from: 1 to: 572

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81 CGAGTCGCGTTGCTTGGCGA.....AGAAATATGTCG 112
||| |||||||||
157 ArgArgTrpValAlaTrpArgCysArgGluLeuSerSerArgCysCysG 173
113 GCATGCGCCCTCGATGAAATCAAGGA..... 140
||| |||
173 yProProGluProSerThrAlaArgThrArgGlyTrpProValA 190
141 .....AGGTGAAGCCGTCAAAAAGCCCAAGT 167
190 rgSerProProValThrArgArgSerSerArgTrpLysArgProPro 206
168 GCTGTTTGAAGACAAAGAAATCGCGGCTAGTATTACTCGCGCGCTT 217
207 .....AsnSerLeuThrCysSerProMe 214
218 CAGGCAAAATCGCGCTATTACCG.....T 243
214 targ.....AlaArgTrpThrProAlaAspGlyAlaCysTrpPheCys 229
244 GG...CGAAAGCGCTACTTCAGTCAGTCGTGATTGCGCTTGAAGCAA 290
229 rpThrArgCysAlaProProSerAlaGlyArgHisLeuProGlyArgSer 245
291 .....CGACGAATCGAGTTGCAACCTACGTACTCTGAAGCGCTGG 331
246 ThrAsnProArgAlaArgCysArgProThrArgLeu..... 258
332 CAAAATTGACGACGAAAGAGTGGCGCAACCTGATTCAATCAGCGTTA 381
259 .....ProAsnAlaPro...ProArgAsnSerArg***C 269
382 TGGACTCGCGTTCGACACCGCTCGGTTTCAGCAA..... 413
269 ysIleCysTrpArgTyrValMetLeuGlnArgArgThrSerCysGlyIle 285
414 .....ATCCCTGCGGTAGATGCCG 433
286 AspSerArgAsnTrpValSerArgTrpProSerProLeuLeuArg..Pro 301
434 AGCGGTTCGCCTTCGTCATCGCATCGATGGACACCAATC...CGCTGGCT 480
302 ThrAlaThrProThrSerThrProThrProValProProTrpIly 318
481 GCGACCCCTACGG.....TCATCATCAAGAAGC 509
318 sProAspTrpArgTrpGlyGluLeuAlaGlySer***SerArgArgSerV 335
510 CGCGCAAGACTTCAAC.....GCGGCC 532
335 alProGlyProAlaAspCysArgProValAlaGlyArgGlyAlaLaPro 351
533 TGTGTGTTATGACCGCTGACCGAAACGTAAATCCATGTTGTAAAGCA 582
352 CysTrpArg...SerSerThrAlaThrValProProSerCysSerProGl 367
583 GCAGGCGCAGACGTCGCGTCTGAAATGCTGCAATATCGAACAACATGA 632

```

367 yArgAlaProAlaCys ..... 372  
633 ATTTGGCGCGCCGATCTCGCGGCTTGAGTGGCAGCAGACATTTCATTTCA 682  
373 .....CysAsp 374  
583 TCGAGCCAGTCGGCGCGCAATAAACCCTGTGGACCATCAATTATCAAGAC 732  
375 ArgValGlnThrPro\*\*\*HisArgPro..... 383  
733 GTGATTGTATCGGACGCTTTGTCGTACACAGCGCTCTGAATACCGAGCG 782  
384 .....ProIleSerValProThrS 390  
783 CGTGGTTG...CCTTTGGCGGCGCTGCAAGTCAACAACCGCGCTCTTGC 829  
390 erTrpCysGlyProTrp\*\*\*Thr...ProAlaProArgThr\*\*\*TrpCys 405  
830 GTACCG.....TTTGGTGCAGAGGTGCTCAACTTACGCGCGCGAA 873  
406 CysProMetAlaMetTrp.....ProProLysAs 415  
874 TTGG.....TTGACGCGGCAACCGCTGATTTCCGTTCCGGTATT 914  
415 nTrpTrpProGlyValProArgArgSerAlaGlyAlaSerThrTrpTyr. 431  
915 GAACGGTCGATTCACAAAGCGCGCATGATTATTGGGACGCTACCACA 964  
432 .....ProCysArgProAspArgTrpCysArgGlyTrp 442  
965 ATCAGATTTCGTTATCGAAGAGCGCGCAGCAAGAGCTTCGCGCT.. 1012  
443 ProArgTrpProCysMetThrArgProAlaArgProSerThrThrAlaTh 459  
1013 .....GGTTGCGCGC..... 1024  
459 rAlaTrpProValProProValLeuProGlyThrAspArgCysAlaLeuP 476  
1025 ..AGCGGACAATACTCCATCAGCGCGCACCACCTCTCGGC..... 1063  
476 roProLysArgArg\*\*\*ProGlyProValProAlaSerArgAlaThrVal 492  
1064 ...ATTTCCTAAAAACAACACTCTCAAGTTCA.....CG 1095  
493 TrpValSerArgAlaThrArgCys\*\*\*SerSerProThrMetSerProAr 509  
1096 ACAGCGCTCAGCGCGCGGACCGCGCCATGTTACGATCGGCACCTATGA 1145  
509 gArgProSerValTrpSerThrCysCysTrpHisArgGluAlaIle.... 524  
1146 GCGCGTAATGCGGTGGACATCTCTACCTGCTTTTGGCGGATTTAA 1195  
525 .....Trp\*\*\*ArgCys..... 528  
1196 TCGTCGCGGATACCGACAGCGCGAGGCTTTGGTT...GCTTGAATTG 1242  
529 ...\*\*\*LeuAlaProAla\*\*\*ProLysThrTrpLeuSerSerTrpAsnG1 544  
1243 GACGAAGAGACCTCGCTTTGTGAGCTTCGCTCCCGCGGCAATACGA 1292  
544 yMetCysThrThrThrIleGlnAlaProSerTrpSerProThrAlaProA 561  
1293 ATACGCGC.....CGCTGT 1306  
561 spHrAlaAlaThrArgCys 567

seq\_name: /sids1/gcgdata/geneseq/geneseq-emb1/AA2001.DAT:ABB64198  
seq\_documentation\_block:  
ID: ABB64198 standard; Protein: 2406 AA.  
XX  
AC ABB64198;

XX 26-MAR-2002 (first entry)  
DT Drosophila melanogaster polypeptide SEQ ID NO 19386.  
XX Drosophila; developmental biology; cell signalling; insecticide;  
KW pharmaceutical.  
KW Drosophila melanogaster.  
OS WO200171042-A2.  
PN 27-SEP-2001.  
XX 23-MAR-2001; 2001WO-US09231.  
XX 23-MAR-2000; 2000US-191637P.  
PR 11-JUL-2000; 2000US-0614150.  
XX (PEKE ) PE CORP NY.  
XX Venter JC, Adams M, Li PWD, Myers EW;  
PI WPI; 2001-656860/75.  
XX N-PSDB; ABL08301.  
DR New isolated nucleic acid detection reagent for detecting 1000 or more  
XX genes from Drosophila and for elucidating cell signalling and cell-cell  
PT interactions -  
XX Disclosure; SEQ ID NO 19386; 2lpp + Sequence Listing; English.  
PS The invention relates to an isolated nucleic acid detection reagent  
XX capable of detecting 1000 or more genes from Drosophila. The invention is  
CC useful in developmental biology and in elucidating cell signalling and  
CC cell-cell interactions in higher eukaryotes for the development of  
CC insecticides, therapeutics and pharmaceutical drugs. The invention  
CC discloses genomic DNA sequences (AB116176-AB130511), expressed DNA  
CC sequences (AB101840-AB116175) and the encoded proteins  
CC (ABB57737-ABB72072).  
CC The sequence data for this patent did not form part of the printed  
CC specification, but was obtained in electronic format directly from WIPO  
CC at ftp.wipo.int/pub/published\_pct\_sequences.  
XX Sequence 2406 AA;  
SQ

alignment\_scores:  
Quality: 108.00 Length: 473  
Ratio: 0.519 Gaps: 28  
Percent Similarity: 43.975 Percent Identity: 22.833  
alignment\_block:  
US-09-303-518D-131/rev x ABB64198 ..  
Align seg 1/1 to: ABB64198 from: 1 to: 2406  
1298 CCGTAT.....TCGTATTCCCGCGCAGCAGGAGCT 1267  
|||||  
1266 ProTyrSerGlyLeuThrHisGlySerTyrLeuProProValLeuProVa 1282  
1266 G.....CACAAAGCGAGGT 1253  
1282 lAlaThrProAsnLeuSerAsnLeuProProThrGlnHisArgSer...S 1298  
1252 CTTCTTCGTCCTCAATCCAAAGCAACCCAAAGCCTCGCGCTGTCGGTATCG 1203  
|||||  
1298 erAspSerArgAsnSerArgGluSerProAla..... 1308  
1202 CCGACGATTAAATCGCGCAAAAGCAAGGTAGG....AGGATGTCCAACGG 1156  
|||||  
1309 ...SerLeuLysSerThrProSerAsnIleGlyLeuAsnValSerMetAl 1324

1155 CATTACG...CGTCATAAGTCCGATCGGT.....ACCA 1124  
1324 aProThrLeuArgSerIleThrProLeuAsnAsnSerSerAlaIleSerS 1341  
1123 TGGCGGGTCCGCGCGTTCACGGCTGTGGT.....AACTGAAG 1083  
1341 erGlyAlaSerGlnProValSerValProSerAlaAsnSerThr 1357  
1082 AGTTTGTGTTTAAAGAAATGGCGAGAGTGGTG.....CGCGTGA 1042  
1358 AlaLeuSerMetSerAsn...ProHisIleSerHisSerHisValPr 1373  
1041 GGAGTATTGTCGGC.....TCGGCGCAACCC 1013  
1373 oAlaTyAlaSerGlyAlaPheSerSerAlaAlaAlaGlyThrSerT 1390  
1012 AGCGCAACAGCTCTTG.....CTCGGCTTCTTCGATAACGGAAATC 969  
1390 hrProAsnSerGlyLeuSerThrLeuAlaValThrSerLeuSerThr... 1405  
968 TGATTGTGTAGGTCCCAATAATCATCGCGCTTGTGCAATCGCAC 919  
1406 .....SerAlaAlaPro.....GlnPr 1411  
918 GTTCAATACCGAACCGAATCAGCGG...TGTCGGCGGTCAACCAATT 872  
1411 oHisSerHisPheProGlnSerThrGlnMetLeuProGlnSerGlyAsn 1428  
871 CGCGGCGGTAAAGTTGAGACACCTTCGCAACCAACAGGTACGCAAG 822  
1428 heSerValSer..... 1432  
821 CGCGGTTGTTGACTTGCAGCGCGCCCAAGCAACCGCGTTCGGTATT 772  
1433 ...HisLeuLeuThrThrHisProMetSerSerGlnAsnGlnProMetVa 1448  
771 CAGCGGCGCTGTACGAACAACGTCGATACCAATCAGCTTGTATAT 722  
1448 larG..... 1449  
721 TGATGTCACACAGGTTTATTTCGCGCGACTGCTGCATGAATGAATG 672  
1450 .....CysGlySerThrLeuTySerGlnSerSerAla..... 1460  
671 TCGCTGCCACTCAACCGCGAGGATCGCGCGCCCAATTCATGTGTT... 624  
1461 .....AlaAlaThrAlaProProSerAlaAlaAla 1471  
623 .....TCGATATTGGCAG 611  
1471 aValSerAsnPheThrProSerValLeuAlaValGlnSerLeuThrTh 1488  
610 CATTTTCAGCGGACAGTCTCGCTGCTGCTTACACACATGATGATTTA 561  
1488 laValThrSerSerSerProSerThrLeuSerSerValIle 1504  
560 CGTTCGGTC.....AGCGGCTCAATACCAACAGGCC 529  
1505 GlnLysValIleSerProLysGlyLysSerProCysAsnLysAspArg 1521  
528 CGGTTTGAAGTCTTCGCGCGCTTCTTTCATGATGATGATGATGATG 479  
1521 pSerSerTySerSerProAlaAsnAlaValThrThrCysAlaPro 1538  
478 CCAGCGGATTGGTTCATCGCATGACGAAGATGGCGAAGCGCTCGCA 429  
1538 hrThrProIleValSer.....SerGlySerAla 1547  
428 TCTACGCGAGGATTTCCTCAACGCGGCGGTGGCAACCGCATCCATA 379  
1548 ArgProThrProProLeuSerAsn..... 1555  
378 GCCTGATTGAATCAGTTGGCGCGCACTTTTCGCTGCTCAATTTGCCA 329

1556 .....CysThrSerMetGlyIleGlyMetVal 1565  
328 GCGCTTACAGTACGTAGCGTTCG.....AACTCGATTTCG...TCG 291  
1565 snAlaAlaSerThrAlaArgSerCysAsnAlaIleSerProLeuSer 1581  
290 TGGCTTCAACGGCAATCAGACTGACTGAAAGTACGGGCTTTTCGCCACG 241  
1582 IleProAlaThr..... 1585  
240 GTGAATACGCGGATTTTTCCTGAGCGCGCAGTAAATACACTACGCCCG 191  
1586 .....AlaGlyIleHisValSerAla.....ThrAsnPro 1596  
190 GATTCTTTTCTCTCAACAGCACTTGGCCTTTTTCGACGGCTTCACCT 141  
1596 erPhe.....GlnSerSerSerTyThrProThrProLeuAlaPro 1610  
140 TCCTTGATTTCATCGAGGGCGCATGCCGACATATTCTTCGCCAAGCAA 91  
1611 Pro.....SerSerProSerPr 1616  
90 CGGACTTCGGTAATGGCC 72  
1616 oAlaThrSerSerAlaAla 1622

seq\_name: /SIDS1/gcgdata/geneseq/geneseq-emb1/AA2001.DAT: AAB59827

seq\_documentation\_block:

ID AAB59827 standard; Protein; 1592 AA.

AC AAB59827;

DT 04-APR-2001 (first entry)

XX Protein #4 encoded by TutD/E gene.

XX Toluene degradation; enzyme; waste degradation; TutE; TutD.

XX Thauera aromatica.

OS Xanthomonas maltophilia.

OS Geobacter metallireducens.

OS Azorarcus toluyticus.

PN WO200072650-A2.

XX 07-DEC-2000.

XX 24-MAY-2000; 2000WO-US14298.

XX 01-JUN-1999; 99US-0323872.

XX (UYOH-) UNIV OHIO.

XX Coschigano PW;

XX WPI; 2001-041080/05.

XX N-PSDB; AAF23627.

XX Composition comprising toluene degrading enzyme useful for biological treatment of organic compounds, especially for degrading toluene or its analogs

XX Disclosure; Fig 12; 122pp; English.

XX The present invention relates to toluene degrading enzyme genes and proteins tutH (see AAF23629 and AAB59831), tutI (AAF23630 and AAB59832), tutF (AAF23631 and AAB59833) and tutG (AAF23632 and AAB59834). The toluene degrading enzymes are homologues of pyruvate formate lyase. The toluene degrading enzymes are useful for biological treatment of organic compounds and in particular for the degradation of toluene and its analogs contained in liquid or solid waste source. The present sequence



```
seq_name: /SIDSL/gc9data/geneseq/geneseq-emb1/AA2001.DAT: AAB27242
seq_documentation_block:
ID   AAB27242 standard; Protein; 571 AA.
XX  AC   AAB27242;
XX  DT   27-MAR-2001 (first entry)
XX  DE   Human EXMAD-20 SEQ ID NO: 20.
XX  KW   Extracellular matrix and adhesion-associated protein; EXMAD; cancer;
XX  KW   inflammation; reproductive disorder; cardiovascular disorder;
XX  KW   immune disorder; musculoskeletal disorder; developmental disorder;
XX  KW   gastrointestinal disorder; cell proliferation disorder.
XX  OS   Homo sapiens.
XX  PN   W0200068380-A2.
XX  PD   16-NOV-2000.
XX  PF   10-MAY-2000; 2000MO-US12811.
XX  PR   11-MAY-1999; 99US-0133643.
XX  PR   23-AUG-1999; 99US-0150409.
XX  PA   (INCY-) INCYTE GENOMICS INC.
XX  PI   Bandman O, Hillman JL, Tang YT, Lal P, Yue H, Baughn MR, Lu DAM;
XX  PI   Azimzai Y;
XX  DR   WPI; 2001-007395/01.
XX  DR   N-PSDB; AAC66909.
XX  PT   Isolated polynucleotide encoding extracellular matrix or
XX  PT   adhesion-associated protein (EXMAD) useful for diagnosing, treating, or
XX  PT   preventing disorders associated with expression of EXMAD such as
XX  PT   proliferative, immune and genetic disorders.
XX  PS   Claim 1; Page 106-107; 129pp; English.
XX  CC   The present invention provides the protein and coding sequences for 25
XX  CC   novel extracellular matrix and adhesion-associated proteins (EXMADs).
XX  CC   These are designated EXMAD-1, EXMAD-2, EXMAD-3, EXMAD-4, EXMAD-5,
XX  CC   EXMAD-6, EXMAD-7, EXMAD-8, EXMAD-9, EXMAD-10, EXMAD-11, EXMAD-12,
XX  CC   EXMAD-13, EXMAD-14, EXMAD-15, EXMAD-16, EXMAD-17, EXMAD-18, EXMAD-19,
XX  CC   EXMAD-20, EXMAD-21, EXMAD-22, EXMAD-23, EXMAD-24 and EXMAD-25. They are
XX  CC   useful in the prevention and treatment of cancers, cell proliferation,
XX  CC   cardiovascular, reproductive, immune, musculoskeletal, developmental and
XX  CC   gastrointestinal disorders and inflammation.
XX  SQ   Sequence 571 AA;

alignment_scores:
Quality: 106.50      Length: 514
Ratio: 0.467         Gaps: 19
Percent Similarity: 44.358      Percent Identity: 18.872

alignment_block:
US-09-303-518D-131/rev x AAB27242
Align seg 1/1 to: AAB27242 from: 1 to: 571

1340 CCTTCCTCTCAATGGTTTCAGACACTTCGCCAACAGCGCGCGATTTC 1291
||| ||| :|||:||||| :|||:||||| :|||:||||| :|||:|||||
23 ProMetMetProThrThrSerGlyThrSerGlnAlaSerSerPheAs 39
1290 GTATTTCCCGGCGACGAGCTGACAAAGCGAGGTCTCTCTCTGTC 1241
:|||||:||||| :|||:||||| :|||:||||| :|||:|||||
39 nThrAlaLysThrSerThrSerLeuHis.....SerHisThrSerS 53
```

```
1240 ATTCCACGACCAACCAAGCCCTGCGCGTGTCTCGGTA..... 1206
:|||||:||||| :|||:||||| :|||:||||| :|||:|||||
53 erThrHisHisProGluValThrProThrSerIleThrAsnIleThrLeu 69
1205 .....TCGCCGACGATTAAATCGCGCAAAAG 1180
:|||||:||||| :|||:||||| :|||:||||| :|||:|||||
70 AsnProThrSerIleGlyThrThrProValAlaHisThrThrSerAl 86
1179 CAAGGTAGGAGGATGCCAACGGCATACGCGCTCATAGTCCGATCG 1130
:|||||:||||| :|||:||||| :|||:||||| :|||:|||||
86 aThrSerSerArgLeuThrThrProPheThrThrHisSerProThrG 103
1129 GTACC.....ATGGCGCGTCCCGCCGTCGCGTGTCTGCGAATTC 1086
|||||:||||| :|||:||||| :|||:||||| :|||:|||||
103 lysSerProIleSerSerThrGlyProMetThrAlaThrSerPheGln 119
1085 AAGAGTTTGTTTTATAGAAATGCCGAGAGTGGTGGCGTGATGGAGTA 1036
:|||||:||||| :|||:||||| :|||:||||| :|||:|||||
120 ThrThrThrTyrThrPro..... 126
1035 TTGTGCGGCTGCGGCGCAACCCAGCGAACAGCTCTTTGTCGCGCCTT 986
:|||||:||||| :|||:||||| :|||:||||| :|||:|||||
127 .....ProSerHisProGlnThrThrLeu..... 134
985 CTTCCGATACGGAATCTGTGTTGTTAGTCCCAATAATCATCGCGG 936
134 ..... 134
935 CCTGTGCAATCGACCGCTTCAATACCGAA.....CCGGAATC... 897
||| :|||:||||| :|||:||||| :|||:||||| :|||:|||||
135 ProThrHisValProPheSerThrSerLeuValThrProSerThrHi 151
896 .....ACGCGGTTGTCGCGCTCAACCAATTCGC 869
|||:||||| :|||:||||| :|||:||||| :|||:|||||
151 sThrValIleThrThrHisThrGlnMetAlaThrSerAlaSerIleH 168
868 CGCGCGTAACTTCAGACACTTCGCCACCAACGCTACGACGAGGCGC 819
|||||:||||| :|||:||||| :|||:||||| :|||:|||||
168 isSerThrProThrGlyThrValProProThrThrLeuLysAlaThr 184
818 GGTTCGTTGCTGACGCGCGCCCAAGCAACGCGCTCGGTATTCAG 769
|||:||||| :|||:||||| :|||:||||| :|||:|||||
185 GlySerThrHisThrAlaProMetThrValThr..... 196
768 ACGCGCTGTACCAACAAACGTCGCTACATCAGCTTCTGATATGTA 719
|||||:||||| :|||:||||| :|||:||||| :|||:|||||
197 .....ThrSerGlyThrSerG 202
718 TGTCTCACACGGTTTATTTCGCGCGACTGCGCTCGATGAAATGAATGTC 669
:|||||:||||| :|||:||||| :|||:||||| :|||:|||||
202 lnThrHisSerSerPheSerThrAlaThrAlaSerSerPheIleSer 218
668 .....GTGCCACTCAAGCCGCGAGGATCGGCC 640
:|||||:||||| :|||:||||| :|||:||||| :|||:|||||
219 SerSerSerTrpSerSerTrpLeuPro...GlnAsnSerSerSerArgPr 234
639 GCCAAATTCATGTGTTTCG..... 621
|||||:||||| :|||:||||| :|||:||||| :|||:|||||
234 oProSerProIleThrThrGlnLeuProHisLeuSerSerAlaThr 251
620 .....ATATTGGCAGCATTTTTCAGACGCGCAGC 594
:|||||:||||| :|||:||||| :|||:||||| :|||:|||||
251 hrProValSerThrThrAsnGlnLeuSerSerPheSerProSerPro 267
593 TCTGCGCCTGCTGCTTACACACATGGATTTACGTTCTGTCAGCGGCT 544
|||||:||||| :|||:||||| :|||:||||| :|||:|||||
268 SerAlaProSerThrValSerSerTyrValProSerSerHisSerPr 284
543 CAATACCAACAGCGCGTTCGAAGTCTTCGCGCGCTTCTTTGATGATGA 494
:|||||:||||| :|||:||||| :|||:||||| :|||:|||||
284 ocGlnThrSerProSerValGlyThrSerSerSerPheValSerAlap 301
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189 aITyrPheGlyArgAsnAlaTyrGlyValGlnAlaAlaHisAlaPhe 205  
394 CGCACCCGTCGTCAGCAAAATCCCTGCGTAGATGCGAGCCGTCGCG 443  
206 PheAspLysProValGluGluLeuThrAlaAlaGluGlyAlaValLeuAl 222  
444 CATCTTCGTC.....AATGCGATGAC.....ACCAATCGCGTG 478  
222 alaSerIleGlnLeuProSerGlnLeuAspProTrpThrAsnProValG 239  
479 CTGCGCAGCCCTACGTCATCATCAAGAGAGCGCGCGAAGACTTCAAAGC 528  
239 luAlaGlu.....ThrArg 243  
529 GGCCTGTGTTGATTGAGCGCGCTGACGCAAGTAAATCCATGTGTAA 578  
244 TrpAsnTyrValMetAspLysLeuValGlu..... 253  
579 ACACAGCGCGAGACGTCGCTGCGAATAATGCTGCCAATATC.....G 622  
254 ....IleGlyAlaIleSerAlaGluGluArgAlaValAlaThrTyrProG 269  
623 AAACACATCAATTTGGCGCGCGCATCTGCGCGCTTGAGTGGCAGCGAC 672  
269 luThrThrAsp.....ProAlaSerAsnSerAla..... 278  
673 ATTCATTTTCATGACGCGCGCGCGAATAAAACCGTGTGGACCATCAA 722  
279 .....TyrThrGluAlaThrGlyThrAsn.....GlyLeuIleLys 290  
723 TTATCAACAGCTGATTGCTATCGGAGCGTTTGTGTAACAGCGCGCTCTGA 772  
290 sAsnGlnValMetAlaGluLeuSerGluLeuGlyLe..... 302  
773 ATACGAGCGCGTGTGCTTGGCGCGCTGCAAGTCAAC..... 813  
303 ..ThrGluAspValGlnThrArgGlyLeuGlnValThrThrIle 318  
814 AAACGCGCGCTCTGCGTACCGTGTGCGGAGGTGTCTCAACTTAC 863  
319 AspProLysThrGlnGluGlyAlaValGluAlaValGlnAsnGlnLeu... 334  
864 CGCGCGCAATTTGTGACGCGGACACCGC.....GTGATTTCGGTT 907  
335 .....AspLeuLeuSerGluAsnAsnArgAlaAlaValValSerIleA 349  
908 CGGTATTGAACGTCGATGCA.....CAAGCGCGCATGATTATTG 951  
349 spProSerAsnGlyAlaValArgAlaTyrTyrGlyGlyGluAspAlaThr 365  
952 GGAGCTACCATCATCAGATTTCGTTATCGAAGAGCGCGCAAGA 1001  
366 Gly...TrpAspPheAlaAsnAlaProLeuGlnThrGlySerPheLeY 381  
1002 GCTGTTCGCGTGGTTCG.....CCGACGCGGACA 1033  
381 silePheGlyLeuAlaAlaLeuGlnGlnGlyIleProLeuSerGlnP 398  
1034 AATACTCCATCAGCGGACCACTCTCGGCCATTTCCTTAAAAACAAACTC 1083  
398 roTyrSerAlaProValThrVal..... 406  
1084 TTCAAGTTCAGCAGCGCTCAAGCGGCGGCGCGCCCATGTACCGAT 1133  
407 .....GlyAspAlaGlnIleGlyAsnVa 414  
1134 CGGC.....ACTATGACGCGGTAAATCGCGTTGG 1162  
414 lGlyGlySerGlyCysGlySerCysSerIleGluGlnAlaLeu...LeuH 430  
1163 ACATCTTCGCTACCTGCTCTGTCG.....GATTTAATCGTGGC 1203  
430 isSerTyrAsnThrSerPheIleArgLeuGlnGlnAspLeuGluAsnGly 446

1204 .....GATACC....GACAGCGCGCAGGCTTGGGTGCTTGGAA..... 1239  
447 SerGlnAspThrAlaAspMetAlaHisAlaLeuGlyIleAlaLysSerLe 463  
1240 .....TTGGACGAGAGAGACCTGCTTGTGAGCT 1270  
463 uProThrIleProGluThrLeuThrGluAsnGlyGluThrProTyrGluG 480  
1271 TCGTCTGCCCGGCAATACGAATACGCGCGCTG...TTGCGCAAGTG 1317  
480 lyIleIleLeuGlyGlnTyrGluSerArgProLeuAspMetAlaSerAla 496  
1318 CTGGAACACCATTTGAGAAGAGGCG 1341  
497 MetAlaThrIleAlaAsnGluGly 504  
seq\_name: /SIDS1/gcycdata/geneseq/geneseq-emb1/AA2001.DAT:ABG21954  
seq\_documentation\_block:  
ID ABG21954 standard; Protein; 1078 AA.  
XX AC ABG21954;  
XX AC  
XX 18-FEB-2002 (first entry)  
XX  
XX Novel human diagnostic protein #21945.  
XX Human; chromosome mapping; gene mapping; gene therapy; forensic;  
KW food supplement; medical imaging; diagnostic; genetic disorder.  
XX Homo sapiens.  
XX  
XX WO200175067-A2.  
XX 11-OCT-2001.  
XX 30-MAR-2001; 2001WO-US08631.  
XX 31-MAR-2000; 2000US-0540217.  
PR 23-AUG-2000; 2000US-0649167.  
XX (HYSE-) HYSEQ INC.  
XX Drmanac RT, Liu C, Tang YT;  
XX  
XX WPI: 2001-639362/73.  
DR N-PSDB; AAS86141.  
XX  
PT New isolated polynucleotide and encoded polypeptides, useful in  
PT diagnostics, forensics, gene mapping, identification of mutations  
PT responsible for genetic disorders or other traits and to assess  
PT biodiversity  
XX Claim 20; SEQ ID No 52313; 103pp; English.  
XX  
XX The invention relates to isolated polynucleotide (I) and  
XX polypeptide (II) sequences. (I) is useful as hybridisation probes,  
XX polymerase chain reaction (PCR) primers, oligomers, and for chromosome  
XX and gene mapping, and in recombinant production of (II). The  
XX polynucleotides are also used in diagnostics as expressed sequence tags  
XX for identifying expressed genes. (I) is useful in gene therapy techniques  
XX to restore normal activity of (II) or to treat disease states involving  
XX (II). (II) is useful for generating antibodies against it, detecting or  
XX quantitating a polypeptide in tissue, as molecular weight markers and as  
XX a food supplement. (II) and its binding partners are useful in medical  
XX imaging of sites expressing (II). (I) and (II) are useful for treating  
XX disorders involving aberrant protein expression or biological activity.  
XX The polypeptide and polynucleotide sequences have applications in  
XX diagnostics, forensics, gene mapping, identification of mutations  
XX responsible for genetic disorders or other traits to assess biodiversity  
XX and to produce other types of data and products dependent on DNA and  
XX amino acid sequences. ABG00010-ABG30377 represent novel human

CC diagnostic amino acid sequences of the invention.  
CC Note: the sequence data for this patent did not appear in the printed  
CC specification, but was obtained in electronic format directly from WIPO  
CC at ftp.wipo.int/pub/published\_pct\_sequences.  
XX  
SQ Sequence 1078 AA;

alignment\_scores:  
Quality: 106.50 Length: 486  
Ratio: 0.461 Gaps: 21  
Percent Similarity: 47.531 Percent Identity: 21.811

alignment\_block:  
US-09-303-518d-131/rev x ABG21954 ..  
Align seg 1/1 to: ABG21954 from: 1 to: 1078

1337 TCCTCTCAATGTTTCCAGCACTTTCGCAACAGC..... 1302  
320 AlaphaLeuValThrSerSerLeuSerGluSerValPheLeuArgAs 336  
1301 .....GGGCGGATTCGATTTCGCGCGGAGACGACGCTGCACAAAG 1259  
336 pValPheGlyLeuSerAlaPheLeu.....ArgSerLeuAlaLys 350  
1258 CGAGGCTCTTCTCGTCCAAATTCACCAAGCAACCCAAAGCTCGCGCTGTCG 1209  
350 eAsnSerLeuGlyLysIleSerPheLeuSerSerLeuLeuAlaSer 366  
1208 GTATCGCGGAGATTAATCG.....CGCAAAAG 1180  
367 AlaserValThrIleSerSerProSerThrSerSerLeuGlnArgLysAl 383  
1179 CAAGGTAGGAGGATG...TCCACGCGATTACCGCTCATAGTGCAG 1133  
383 aSerPheAlaArgLeuProSerSerGlyLeu.....Pros 395  
1132 TCGGTACCATCGCGCGCGCGCTGAGCGTGTGCTGCTGAACTGAAG 1083  
395 eGluThrCysIleArgLeuAlaSerAlaSerProValLysGlyLeuLys 411  
1082 AGTTTGTGTTTATAGGAATGCCGAGAGTGTGCGCGCTGATGAGTATT 1033  
412 ArgVal.....SerValLeuIleThrSerPheLe 421  
1032 GTCCGGCTGCGCGCAACCCAGCGACAGCTCTTTGCTCGCGCTCTT 983  
421 uSerLeuAlaGlySerSerProProArgIleAlaSerSerGluProSer 438  
982 CGATAACGGAATCTGATTGTTGGTAGCTCCCAATAATCATGCGCGCT 933  
438 eLeuPheASP.....AlaVal 443  
932 TGTGCAATCGCACCGTTCAATPACCGAACCAGGAAATCAGCGGTTGTCGC 883  
444 SerSerMetGlyProValSerSerArg.....SerMetAspAl 456  
882 GTCAACCAATTCGCGCGGCTAAGTTGAGACACCTTCGCCACCCAAACGG 833  
456 aSerSerLeuSerProThrSer..... 464  
832 TAGCGAAGCGCGGTTGTTGACTTTCAGCGCGCCCAAGGCAACACG 783  
465 .....LeuThrAlaThrProProSerAlaSerPro 474  
782 CGCTCG.....GTATTCAGAGCGCTTTCAGCAACAACTCCGAT 742  
475 SerSerAspValAlaAlaSerLeuArgAlaAlaThrSer\*\*\*ArgCysSe 491  
741 ACCAATCAGCTTCGATTAATGATGGTCCACACGCTTTTATTCGCGCGA 692  
491 rLysPheArgAla.....AlaAlaSerPheAlaSerT 502

691 CTGGCTCGATGAATGATGTGCGTGCCTCAAGCGCGCAGGATCGGG 642  
502 hAlaPheSerLys.....ProCysSerProSerGluAlaAla 514  
641 ...CGGCCAAATTCATGTTTCGATATTCGACAGATTTTCAGACGCGAC 595  
515 LeuSerValSerSerThrValSerIleLysAlaCysPheSerGlyAlaSe 531  
594 GTCTCGCGCTGCTGCTTTACACATGATTTTACGTTTCGCTCAGCGCGC 545  
531 rSer.....LeuLeuArgProThrAlaSerP 540  
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540 hAlaAlaAlaCysProMetGlySerAspProProSerProSerLeuLeu 556  
494 ACCGTAGGTCGGCAGCCAGC.....GG 472  
557 ThrProSerAlaSerAlaAlaProSerAlaProLeuSerMetProGlyGl 573  
471 ATTGCTGCCATCGCATTCGACGAGATGGCAAGCGCTCGGCATCTACGG 422  
573 yIleSerLeuPheSerValAsnSerSerAlaAsnSerGlyCysSerAlaV 590  
421 CAGGGATT...TTGCTGAACGACGCGGTGCGAAGCGCAGTCCATTAAGCT 375  
590 alGlyPheHisAlaLeuLysGlyAlaSer\*\*\*ThrAlaProAlaSerPro 606  
374 GATTGAATCAGGTTGCGCGCACCTTTTCGCTGCTCAATTTTCCGACGCGC 325  
607 His.....SerAlaAlaSerProPheProCysThrMetSerAlaSerLe 621  
324 TTCAGGTAGCTAGCGTTCGAACTCGATTTCGCTGCTTCCTCAACGGCAA 275  
621 uCysTrpGlyAlaSerSerSerIleAlaSerPhePheSerSerValT 638  
274 TCACGACTGACTGAAGTACGCGCTTTTCGCCACGCGTGAATACGCGCGATT 225  
638 yrVal.....PheCysThrAsnCysTrpAspSerLeu 648  
224 TTCCTGTAAGCGCGCGCTAAATACTACGCCCGGATTTCTTTTG..... 180  
649 PheGluLeuIleAlaAlaValSerSerThrMetLysPheIleValGlyAl 565  
179 .....TCTTCAACAGCAGCTTGGCTTTTTCACGCGCTT 146  
665 aProArgAlaAArgThr\*\*\*AlaAsnGlyAsnGluSerPheLeuProLeuV 682  
145 CACCTTCTCTGATTTTCATCGAGGGCGCATGCCACATATTCT..... 102  
682 aIserThrSerSerPheLeuGlyGlyPheProProSerSerSerLeuLeu 698  
101 TCGCCCAAGCAACGCGACT.....TC 82  
699 ThrProAspSerAlaSerGlnTrpGlnTrp\*\*\*ProAsnAlaLysGlySe 715  
81 GGTATGCGCGCGCGCTGATAAATGACTTTCGCTCGCGTTCGCCGCGATGG 32  
715 rValIleIleCysProAlaProSerSerAsnProGlyArgProAlaPheL 732  
31 GCAGATT 24  
732 euSerPhe 734

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seq\_documentation\_block:  
ID AAW87504 standard; Protein; 1061 AA.  
XX  
AAW87504;  
XX  
23-FEB-1999 (first entry)  
DT





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914 oProGlySerThrAspSerThrSerAlaCysThrProThrProThrC 931
1018 GCGCCGAGCGGACAAATACTCC.....ATCAGCG 1049
1031 yHisCysAlaGlyGlyLeu.SerValLeuThrPheHisProValThrAl 947
1050 CACACACTCGGCGCATTCCTAAACAAACACTCTTC..... 1086
1064 aThrAlaProGlySerProAlaProGlyGlyLeuTrpGlyThrAlaAlaG 964
1087 .....AAGTCAGACACCGCTCAACGGCGGCGCGC 1119
1094 lyLeuTrpGlyTrpAlaGlnAlaThrValGlyAspTrpThrArg 980
1120 GCCATGTCAGCGGCGCACTTATGACGCGTAATGCCGTTGGACATCCT 1169
1181 SerAlaValProValGlyArgLysAlaSer..... 990
1170 GCCTACCTTGTCTTTCGCGGATTTAATCGTCGGCATFACCGACGCGC 1219
1191 .....ArgaspProAlaProGlyAsp..... 997
1220 AGGCTTTGGTGTCTTGAATTCGACGAAGACCTCGCTTTGTCAGC 1269
1298 .....Gly.SerProValTrpSerGlnLysCysGluLeuSerAlaTh 1011
1270 TTCCTCTGCGCGGCAATACGAATACGCGCGCT.....GTTGCG 1310
1011 rGlnAla.ProSerGlnLeuAspSerLeuProAlaThrValArgValLys 1027
1311 CAAAGTGCTGGA 1322
1028 ArgGlnAlaGly 1031

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seq\_name: /SIDSI/gcgdata/geneseq/geneseq-emb1/AA1999.DAT:AAW87503

seq\_documentation\_block:

XX AA87503 standard; Protein; 1212 AA.

XX AC AA87503;

XX DT 23-FEB-1999 (first entry)

XX DE Human N-methyl-D-aspartate receptor subunit encoded by clone NMDA22.

XX KW Human; N-methyl-D-aspartate receptor; NMDAR2C;

XX KW NMDA-activated cation-selective ion channel; glutamate receptor.

XX OS Homo sapiens.

XX PN US5849895-A.

XX PD 15-DEC-1998.

XX PF 20-APR-1994; 94US-02311193.

XX PR 20-APR-1994; 94US-02311193.

XX PR 20-APR-1993; 93US-0052449.

XX PA (SIBI-) SIBIA NEUROSCIENCES INC.

XX PI Daggett LP, Lu C;

XX DR WPI; 1999-069812/06.

XX DR N-PSDB; AAV82909.

XX PT DNA encoding N-methyl-D-aspartate receptor subunit - useful for the

XX PT assembly of functional glutamate receptor subunits

XX PS Example 3; Columns 253-262; 203pp; English.

XX PS The present sequence represents a human N-methyl-D-aspartate (NMDA)

CC

receptor subunit (NMDAR). The nucleic acid sequence does not contain the 366 5'-most nucleotides, by the insertion of 11 nucleotides between nucleotides 1300 and 1301, nor the 15 nucleotides at positions 1960-1974, nor the 1061 3' nucleotides, as set forth in AAV82889. The cDNA sequence is derived from clone NMDA21. The NMDAR subunits contribute to the formation of NMDA-activated cation-selective ion channels. In addition to being useful for the production of NMDA receptor subunit proteins, the nucleic acids are also useful as probes to identify and isolate nucleic acids encoding related receptor subunits. Functional glutamate receptors can be assembled from several NMDA receptor subunit proteins of one type (homomeric) or from combinations of subunit proteins of different types (heteromeric). The present invention also comprises methods for using such receptor subunits to identify and characterize compounds which affect the function of such receptors, e.g. agonists, antagonists and modulators of glutamate receptor function. The invention also comprises methods for determining whether unknown protein(s) are functional as NMDA receptor subunits.

XX Sequence 1212 AA;

alignment\_scores:

Quality: 106.00 Length: 490

Ratio: 0.613 Gaps: 28

Percent Similarity: 35.306 Percent Identity: 23.265

alignment\_block:

US-09-303-518D-131 x AA87503 ..

Align seg 1/1 to: AA87503 from: 1 to: 1212

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840 LeuAlaProSerArgIleGlyValAlaAlaValArgProHisArgPr 856
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76 ATTACCGAAGTCGCGTGTCTGGCGAAGAATATGTCGCGATCGCGCCCTC 125
   :::: :::: :::: :::: :::: :::: ::::
856' OProAlaArgProArgGlyLeuAlaPro...AlaHisAlaCysProPro. 871
   :::: :::: :::: :::: :::: :::: ::::
126 GATGAAATCAAGGAGGTGAAGCGGTCAAAAAGG...CCAAGTGTGTGT 172
   :::: :::: :::: :::: :::: :::: ::::
872 .....ProThrArgPro.GlnSerArgAlaProArgAlaG 883
   :::: :::: :::: :::: :::: :::: ::::
173 TTGACAGACAAAAGATCGCGGTAGTATTACTGCGCGCGCTTCAGC 222
   :::: :::: :::: :::: :::: :::: ::::
883 yAspArgGlnThrGlyValAlaArgLeu...CysAlaGlyLeuArg. 898
   :::: :::: :::: :::: :::: :::: ::::
223 AAAATCGCGCTATTACCGTGGCGGAAAGCGCGT...ACTTCAGTCAGT 269
   :::: :::: :::: :::: :::: :::: ::::
899 ..SerProArgAlaAlaProArgArgGlyArgProCysProThrSer 914
   :::: :::: :::: :::: :::: :::: ::::
270 CGTGATTCGCGTTGAGGCAACGACCAATCGAGTTCGAACGCTACGTAC 319
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915 ProGluCysArgAlaAlaGlnProGlyArgArgGlyArgCysGlyPr 931
   :::: :::: :::: :::: :::: :::: ::::
320 CTGAAGCGCTGGCAAAATTTGACGACGCAAAAGTGGCGCGCAACCTGATT 369
   :::: :::: :::: :::: :::: :::: ::::
931 oGlyThrAlaGlyGlyThrSerArgProSerGlyPro..... 944
   :::: :::: :::: :::: :::: :::: ::::
370 CAATCAGGCTTATGACTCGCTTCGCCCGCTCCGTCAGCAAAATCCC 419
   :::: :::: :::: :::: :::: :::: ::::
944 ..... 944
420 TGCGGTAGATGCGGACGCGGTTCGCCATCTTGTCAATGCGATGGACACCA 469
   :::: :::: :::: :::: :::: :::: ::::
944 ..... 944
470 ATCCGCTGGTGGCGGACCGCTACGTCATCATCAAGAACGCCCGCGAGAC 519
   :::: :::: :::: :::: :::: :::: ::::
945 .....CysArgPro.....ArgAlaV 950
   :::: :::: :::: :::: :::: :::: ::::
520 TTCAACGCGGCGCTGTGGTATTGATGACCGCGCTGACCAACGTAATCCA 569
   :::: :::: :::: :::: :::: :::: ::::

```



1313 eserPheGlyPheGlyGlnSerAsnGlnGlyLysAspValAlaAspSerL 1330  
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1243 CCAATTCACCAACCAACCAACCTGCGGCTGTCGCTATCCGCGCAGGATT 1194  
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1330 ysLysThrGluAlaProLysThrPheMetPheGlyValSer 1343  
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1193 AAATCGCCCAACCAACCAACCAAGTAGGATGTCACACGGCATTCGCGCTC 1144  
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1344 LysValGluGluProLysThrValSerPheGlyThrGlyIleLysGluTh 1360  
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1143 ATAAGTCGCGATCGATCCATCGCGCTGCGCGCTGACGGCTGTCG 1094  
||| :||||: |||||:||||: |||||:||||: |||||:||||: 1094  
1360 rThAlaThrSerSerThrGluAlaThrAlaProThrProAlaAlaA 1377  
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1093 TGAATTCGAAGAGTTGTTTATAGGAATGGCGAGAGTGGCGCGCTG 1044  
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1377 laProAlaProValGlnPheValPheLysAlaProThrThrAlaThrThr 1393  
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1043 ATGGAGTATTGTC. 1029  
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1394 AlaSerSerLeuThrThrThrIleSerThrThrSerAsnAlaProAlaLe 1410  
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1028 ...GCCTGCGCGCAACCCAGCGCAACAGCTCTTGTGCGGCTTCTT 983  
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1410 uGlyGlyPheSerPheGlyAlaProSerSerSerThrValSerSers 1427  
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982 CGATAACGGAATCTGATTGTTAGTCCCAATATCATCGCGCT 933  
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1427 erThrThrSerThr 1435  
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932 TGTGCAATCGCACCGTTC. 915  
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1436 AlaAlaValLysProMetPheSerTrpSerGlyAlaGlySerAlaValSe 1452  
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914 .....AATACGCAACG. 903  
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1452 rSerThrSerSerGlnGlnProValAlaLysAlaProThrLeuGlyP 1469  
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1469 heGlyValSerSerThrValThrThrThrThrThrThrThrThrThr 1485  
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872 .....TCGCGCGCGTAACTGAGACACCTTCGCA. 843  
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1486 PheAlaPheThrProAlaSerGlyLeuAspProAlaAlaAlaThrSerAl 1502  
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842 .....CCCAAAA. 836  
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1502 aProAlaAlaGlyAlaGlyPheSerPheGlySerGlnSerLysProAlaT 1519  
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835 CGGTACGC. 810  
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1519 hrThrGlnAsnThrGlyThrPhePheGlyGlnProThrAlaValAla 1535  
:||||: |||||:||||: |||||:||||: |||||:||||: |||||:||||: 1535  
809 ACTTCAGCGCCCAACCAACCAACCGCTCGGTATTCAGACGCGCTGT 760  
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1536 ProAlaThrProThrAsnProSerValSerIlePheGlyAlaProAl 1552  
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759 TAGCAACAACCTCGCATAGCAATCAGCTTGATAATTGATGGTCCACA 710  
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674 ATGTGCTGCCACTACAGCGCGCAGGATCGCGCGCCCAATTCATGTGT 625  
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1681 ThrThrProAlaGlyGlyAsnAlaAlaLeuThrGlyLeuPheGlyAsnVa 1697  
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177 TTCAAACAGC. 155  
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1697 lGlyAsnSerLeuAlaGlyValGlyAlaProValAlaAlaThrThrProAlaA 1714  
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154 TGACGCTTCACCTTCCTTGAATTTTCATCAGGCGCGC. 117  
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1714 laThrAlaAlaAlaProLeuThrAsnIlePheGlyAsnProThrProVal 1730  
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116 .....ATGCGGACATATCTTCG. 88  
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1731 AlaAlaAlaAlaProValPheGlySerGlySerThrIleProSerAlaGl 1747  
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87 GACTTCGCTAATGCGCGCGCTCATAAATGACTTCTCGGTCGCCG 38  
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1747 yPheGlyAlaProAlaAlaAlaAlaProLeuAlaAlaProAlaLeuProG 1764  
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37 CGATGGCGCAGATT 24  
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1764 lyAlaPheAsnPhe 1768  
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seq\_name: /SIDS1/gcgdata/geneseq/geneseq-emb1/AA2001.DAT:AAU48589  
seq\_documentation\_block:  
ID AAU48589 standard; Protein; 372 AA.  
XX  
AC AAU48589;  
XX  
DT 27-FEB-2002 (first entry)  
XX  
DE Propionibacterium acnes immunogenic protein #9485.  
XX  
KW SAPHO syndrome; synovitis; acne; pustulosis; hypervellitis;  
KW uveitis; endophthalmitis; bone; joint; central nervous system; ELISA;  
KW inflammatory lesion; acne vulgaris; enzyme linked immunosorbent assay;  
KW dermatological; osteopathic; neuroprotectant.  
XX  
OS Propionibacterium acnes.  
XX  
PN WO200181581-A2.



|||||:|||||:|||||  
321 erProThrMetSerThrArg 327

seq\_name: /SIDS1/gcgdata/geneseq/geneseq-emb1/AA2001.DAT:AAU39645

seq\_documentation\_block:  
ID\_ AAU39645 standard; Protein; 338 AA.

XX AC AAU39645;

XX DT 13-FEB-2002 (first entry)

XX DE Propionibacterium acnes immunogenic protein #541.

XX KW SAPHO syndrome; synovitis; acne; pustulosis; hypertosis; osteomyelitis;  
KW uveitis; endophthalmitis; bone; joint; central nervous system; ELISA;  
KW inflammatory lesion; acne vulgaris; enzyme linked immunosorbent assay;  
KW dermatological; osteopathic; neuroprotectant.

XX OS Propionibacterium acnes.

XX PN W0200181581-A2.

XX PD 01-NOV-2001.

XX PF 20-APR-2001; 2001WO-US12865.

XX PR 21-APR-2000; 2000US-199047P.

XX PR 02-JUN-2000; 2000US-208841P.

XX PR 07-JUL-2000; 2000US-216747P.

XX PA (CORI-) CORIXA CORP.

XX PI Sheiky YAW, Persing DH, Mitcham JL, Wang SS, Bhatia A;

XX PI L'Maisonneuve J, Zhang Y, Jen S, Carter D;

XX DR WPI; 2001-616774/71.

XX DR N-PSDB; AAS59508.

XX PT Propionibacterium acnes polypeptides and nucleic acids useful for  
PT vaccinating against and diagnosing infections, especially useful for  
PT treating acne vulgaris -

XX PS Example 1; SEQ ID No 840; 1069pp; English.

XX CC Sequences AAU39105-AAU68017 represent propionibacterium acnes immunogenic  
CC polypeptides. The proteins and their associated DNA sequences are used in  
CC the treatment, prevention and diagnosis of medical conditions caused by  
CC P. acnes. The disorders include SAPHO syndrome (synovitis, acne,  
CC pustulosis, hypertosis and osteomyelitis), uveitis and endophthalmitis.  
CC P. acnes is also involved in infections of bone, joints and the central  
CC nervous system, however it is particularly involved in the inflammatory  
CC lesions associated with acne vulgaris. A method for detecting the  
CC presence or absence of P. acnes in a patient comprises contacting a  
CC sample with a binding agent that binds to the proteins of the invention  
CC and determining the amount of bound protein in the sample. The  
CC polypeptides may be used as antigens in the production of antibodies  
CC specific for P. acnes proteins. These antibodies can be used to  
CC downregulate expression and activity of P. acnes polypeptides and  
CC therefore treat P. acnes infections. The antibodies may also be used as  
CC diagnostic agents for determining P. acnes presence, for example, by  
CC enzyme linked immunosorbent assay (ELISA).  
CC Note: The sequence data for this patent did not form part of the printed  
CC specification, but was obtained in electronic format directly from WIPO  
CC at ftp.wipo.int/pub/published\_pct\_sequences.

XX SQ Sequence 338 AA;

alignment\_scores:  
Quality: 105.00 Length: 379  
Ratio: 0.695 Gaps: 18  
Percent Similarity: 39.842 Percent Identity: 22.427

alignment\_block:  
US-09-303-518D-131/rev x AAU39645

Align seg 1/1 to: AAU39645 from: 1 to: 338

1031 TCCGGCTGCGGCAACCCAGCCGCTCTTTGGTGGCGGCTTCTTC 982  
:::|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||  
17 ThrGlyCysAlaGlyLeuThrSerLeuSerThrThrArgAlaSerHi 33  
981 GATAACGGAAATCTGATTGTGTAGCTCCCAATAATCATCATCGCGCTT 932  
|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||  
33 sSer.....SerCysValProV 39  
931 GTGCAATCGCACCGTTCAATACCGAAGCAACCGGTTGTTCGCG 882  
|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||  
39 al.....ThrSerProGlySerSerSerLeuSerSer 49  
881 TCAACCAATTCGCGGCGGTAAAGTTGAGACCTTCGCACCCCAACAGGT 832  
|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||  
50 ProThrThrSerAlaIleAlaArgArgLeuArgProArgSerPr 66  
831 ACCGACAGCGCGGTTTGTGCTGACTTGC..... 804  
:::|||||:|||||:|||||:|||||:|||||:|||||:|||||  
66 oThrGln.....LeuThrCysSerGlyTrpProArgArgSerI 79  
803 .....AGCGCGCCCAAGCAACCCAGCGC..... 780  
|||:|||||:|||||:|||||:|||||:|||||:|||||  
79 leProAlaAlaProAlaProThrSerProThrThrArgAlaArgAsnThr 95  
779 .....TCGGTATTACAGCGGCTGTACGAACAACGTCGATAGC 739  
|||||:|||||:|||||:|||||:|||||:|||||:|||||  
96 ProProGlnSerAlaThrArgArgPheProThrSerLeuTrpProlleAr 112  
738 AATCAGCTCTTGATATAATTGATGTCACACAGGTTTATTTCGCGCGGACTG 689  
|||||:|||||:|||||:|||||:|||||:|||||:|||||  
112 gThrThrSer..... 115  
688 GCTCGATGAATAAGTATGCTGCGCCTCAAGCCGCGCAGGA..... 648  
|||||:|||||:|||||:|||||:|||||:|||||:|||||  
116 .....ArgCysLeuAlaThrAlaArgProGlyTrpSerIle 127  
647 .....TCGCGGCGCGCCAAATTCATGCTGTTTC 622  
128 SerThrThrSerSerTrpCysArgValGlyProSerSerAlaCysValTh 144  
621 GATATTGGCA.....GCATTTT 605  
|||||:|||||:|||||:|||||:|||||:|||||:|||||  
144 rAlaLeuSerAlaGlyIleThrValThrThrArgSerGluCysAlaLeuT 161  
604 CAGACGGCAGCTCTGCGCTGCTGCTTAA...CACACATGGATTTCGT 558  
:::|||||:|||||:|||||:|||||:|||||:|||||:|||||  
161 hrProSerThrSerProProAlaThrGlyPheThrTrp..... 174  
557 TCGGTCAGCGGCTCAATACCAACAGCGCGTTTGAAGTCTTCGGCGGC 508  
:::|||||:|||||:|||||:|||||:|||||:|||||:|||||  
175 .....Se 175  
507 TTCTTTTATGATGACCGTAGGTCGGCAGCCAGCGGATGTTGTCATCG 458  
:::|||||:|||||:|||||:|||||:|||||:|||||:|||||  
175 rThrThrMetLeuThrThrSerGlyThrSerArgAlaLeuThrAlaIleA 192  
457 CATTCAGAGATGGGCAACCGCTCGGCATCTACGCGAGGATTTGCTG 408  
:::|||||:|||||:|||||:|||||:|||||:|||||:|||||  
192 rgArgLeuTyrlle.....SerAlaAlaArgTrpProValValgly 205  
407 AACGACGGGTGCGAGCGCAGTCCCAATAGCCTGAT..... 372  
:::|||||:|||||:|||||:|||||:|||||:|||||:|||||  
206 SerArgArgPheProArgProTrpArgThrProSerCysGlyAlaMetPr 222  
371 .TGAATCAGGTTCGGCGGCACCTTTTTCGCTGCTCAATTTTCCGAGCGCTT 323  
:::|||||:|||||:|||||:|||||:|||||:|||||:|||||  
222 oThrLeuArgIleSerArgProThrTrpMetProSerMetThrSerCysI 239

```

322 CAGGTACCTGAGCTTCGAACTCGATTGCTGCTGCTTCA...ACGGCA 276
||||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
239 leGlyThrThrArgArgGly.....SerProIleProSerLeuThrAla 253
275 ATCAGCAGTACTGAGTACGGCTTTTCGCCACGGTGAATA..... 234
||| ||| ||| ||| ||| ||| ||| ||| ||| |||
254 SerThrIleValArgSerArgArgSerValArgTrpValArgSerSe 270
233 ...CGGGGATTTTCCTGAGCGCGCAGTAAATACCTACGCCGAT 188
||||| ||| ||| ||| ||| ||| ||| ||| ||| |||
270 rSerSerSerProHisProArgThrSerProThrGlyThrIleGlys 287
187 TC.....TTTGTGCTTCAACAGCAGTTCGCCCTTTTTCGACGGCTTCA 144
||| ||| ||| ||| ||| ||| ||| ||| ||| |||
287 exThrArgPheArgProSerAlaArgSerTrp..... 297
143 CCTTCCTTGATTTTCATCGAGGGCGCATGCCGACATATTTTCGCCCAAG 94
||||| ||| ||| ||| ||| ||| ||| ||| ||| |||
298 .....SerSerProse 301
93 CAACGCGACTTCG.....GTAATGCGGG 69
||||| ||| ||| ||| ||| ||| ||| ||| ||| |||
301 rSerAlaThrThrSerArgAsnGlyValMetThrGly 313

```

seq\_name: /SIDS1/gcgdata/geneseq/geneseq-emb1/AA1994.DAT:AA44890

seq\_documentation\_block:

ID AAR44890 standard; Protein; 532 AA.

XX AC AAR44890;

XX DT 22-JUN-1994 (first entry)

XX DE Diphtheria toxin delta-142-147-148 mutant.

XX KW DT; protein exotoxin; NAD-dependent ADP-ribosyltransferase; vaccine;  
 KW diphtheria toxin; deletion mutant; mutagen; variant; double mutant;  
 XX reversion mutation; site-directed mutagenesis.

XX OS Corynebacterium diphtheriae.

XX FH Key Location/Qualifiers

FT Protein

1..532

FT /note= "Diphtheria toxin mutant; Glu(142), Val(147)  
 and Glu(148) have been deleted"

FT WO9325210-A.

XX PN 23-DEC-1993.

XX PF 17-MAY-1993; 93WO-US04606.

XX PR 18-JUN-1992; 92US-0901712.

XX PA (HARD ) HARVARD COLLEGE.

XX PI Collier RJ, Killeen K, Mekalanos J;

XX WPI; 1994-007178/01.

XX N-PSDB; AAQ54338.

XX New DNA encoding diphtheria toxin deletion mutants - with no  
 toxicity and low risk of reversion, and derived toxoids and  
 transformed cells, useful in vaccines

XX Claim 11; 42pp; English.

XX Oligonucleotide-directed mutagenesis of the wild-type diphtheria  
 CC gene (specifically the region encoding the DT-A fragment) results  
 CC in deletion of the codons for Val-147 and active site residue  
 CC Glu-148. The resulting mutagen is not toxic, making it useful in  
 CC diphtheria vaccines. The risk of reversion to toxicity is much

CC lower for the 147-148 double mutant than for the prior art 148  
 CC single mutant, while its immunogenicity is not impaired. The  
 CC 147-148 mutant opt. has other amino acid residues subst. or  
 CC deleted, e.g. wild-type Glu(142) is deleted. The specification  
 CC includes the wild-type DT amino acid sequence (see AAR44888) but does  
 CC not include any mutant sequences; the wild-type sequence was modified  
 CC according to the description in the claims to give AAR44890.  
 XX  
 SQ Sequence 532 AA;

alignment\_scores:

Quality: 105.00 Length: 461  
 Ratio: 0.536 Gaps: 25  
 Percent Similarity: 42.516 Percent Identity: 20.390

alignment\_block:

US-09-303-518D-131 x AAR44890 ..

Align seg 1/1 to: AAR44890 from: 1 to: 532

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31 CCCATCGCGGCGAGACCGGAGCAAGTCATT...TATGACGCGCGCGCCAT 77
||||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
72 ProLeuSerGlyLysAlaGlyGlyValLysValThrTyrProGlyLe 88
78 TACCGAAGTCGCGCTGCTTGGCGCAAGATATGTCGCATGCGCGCCCTCGA 127
||||| ||| ||| ||| ||| ||| ||| ||| ||| |||
88 uThrLysValLeu.....AlaL 94
128 TGAATAATCAAGGAAGGTGAAGCGGTCAAAAAA..... 159
||||| ||| ||| ||| ||| ||| ||| ||| ||| |||
94 euLysValAspAsnAlaGluThrIleLysLysGluLeuGlyLeuSerLeu 110
160 .....GCCAAGTGCTGTTTGAAGACAAAAA 185
||||| ||| ||| ||| ||| ||| ||| ||| ||| |||
111 ThrGluProLeuMetGluGlnValGlyThrGluGluPheIleLysArgPh 127
186 GAATCGCGGCG.....GTAGTATTACTGCGCGGCTTCAGGCAAAA 226
||||| ||| ||| ||| ||| ||| ||| ||| ||| |||
127 eGlyAspGlyAlaSerArgValValLeuSerLeuProPheAlaGlySers 144
227 TCGCGCTATTCAACCGTGGCGGAAAGCGGTACTTTCAGTCAGTCGCGATT 276
||||| ||| ||| ||| ||| ||| ||| ||| ||| |||
144 exSerTyrIleAsnAsnTrpGluGlnAlaLysAlaLeuSerValGluLeu 160
277 GCGGTGAAGGCAACGACCAATCGAGTTCGAA..... 309
||||| ||| ||| ||| ||| ||| ||| ||| ||| |||
161 .....GluIleAsnPheGluThrArgGlyLysArgL 171
310 .....CGCTACGTACCTGAAGCGCTGGCAAAATGAGCA 343
||||| ||| ||| ||| ||| ||| ||| ||| ||| |||
171 yGlnAspAlaMetTyrGluTyrMetAlaGlnAlaCysAla.....G 185
344 GCGAAAGTGGCGGCAACCTGATTCATCAGGCTTA..... 381
||||| ||| ||| ||| ||| ||| ||| ||| ||| |||
185 lyAsnArgValArgArgSer...ValGlySerSerLeuSerCysIleAsn 200
382 .....TGGACTGCGCTTCGACCGCTCCGTTTCAGCAAAATCCCTGCCGT 425
||||| ||| ||| ||| ||| ||| ||| ||| ||| |||
201 LeuAspTrpAspValIleArgAspLysThrLysThrLysIleGluSerLe 217
426 AGATCGCGGAGCGGTTTCGTCATCGATGCGATGGACACCAATCCGC 475
||||| ||| ||| ||| ||| ||| ||| ||| ||| |||
217 u.....LysGluHisGlyProIleLysAsnLysMetSerGluSerProA 232
476 TGGCTGCCGACCCCTACGGTCATCATCAAGAGCGCGCGCAAGACTTCAA 525
||||| ||| ||| ||| ||| ||| ||| ||| ||| |||
232 snLysThrValSerGluGluLysAlaLysGlnTyrLeuGluGluPheHis 248
526 CGCGCGCTGTTGCTGATTGAGCGCGCTGACCGACGTAATCCATGCTGTG 575
||||| ||| ||| ||| ||| ||| ||| ||| ||| |||
249 GlnThrAlaLeuGluHisProGluLeuSerGluLeuLysThr..... 262

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```
576 TAAAGCAGCAGCGCAGACGTCGCGTCTGAAATGCTGCAATATCGAAA 625
262 .....
626 CACATGAATTGGCGCGCCATCTCCGGCTTGAGTGGCAGCAGCATT 675
263 .....
676 CATTTTCATGAGCAGTCGCGGGAATATAACCGTGTGGACCATCAATTA 725
269 ValPhe.....AlaGlyAlaAsnTyrAlaAlaTrrpAlaValAsnVa 282
726 TCAAGACGTGATT..... 738
282 lAlaGlnValIleAspSerGluThrAlaAspAsnLeuGluLysThrThra 299
739 .....GCTATCGGAGCTTG.....TTCGTAACA 762
299 laAlaLeuSerIleLeuProGlyIleGlySerValMetGlyIleAlaAsp 315
763 GCGCGTCTG.....AATACCGAGCGGTGGTTCCTGGCGCGCTGCA 806
316 GlyAlaValHisAsnThrGluGluIleValAla..... 327
807 AGTCAACAACCGCGCTCTTGGTACCGTGTGGTGGCGAAGTGTCTC 856
328 .....GlnSerIleAlaLeuSerSerLeuMetValAlaGlnAlaIleP 342
857 AACTTACCGCGCGGAATTGGTTCACCGGACACCGCGTGAATTCGGT 906
342 roLeu...ValGlyLeuValAsp..... 349
907 TCGGTATTGAACCGTGGCATGCAACAGCGCGCATGATTATTGGGACG 956
350 .....IleGlyPheAlaAlaTyrAsnPheValGluSe 360
957 CTACACAATCAGATTCCGTTATTCGAAGAGCGCGCAGCAAGAGCTGT 1006
360 rIleIleAsnLeuPheGlnValHisAsnSerTyrAsnArgProAlaT 377
1007 TCGGCTGGTGGCGCG.....CAGCG.....GACAAA 1035
377 yr.....SerProGlyHisLysThrGlnPropheLeuHisAspGly 390
1036 TACTCCATFACGCGCAGCATTCTC..... 1059
391 TyrAlaValSerTrrpAsnThrValGluAspSerIleIleArgThrGlyPh 407
1060 .....GGCCATTTCCTAAACAACTCTTCAAGTTCACGA 1096
407 eGlnGlyGluSerGlyHisAspIleLys.....IleT 418
1097 CAGCGCTCAACGGCGCGCGCATGTTACCGATCGGCGCACTTATGAG 1146
418 hrAlaGluAsn.....ThrProLeuProIleAlaGlyVal... 429
1147 GCGGTAATCGCGTGGACATCTGCTACCTG 1179
430 .....LeuLeuProThrIle 434
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seq\_name: /STD1/gcdata/geneseq/geneseq-emb1/AA2001.DAT:ABG21970

seq\_documentation\_block:

ID ABG21970 standard; Protein; 575 AA.

XX AC ABG21970;

XX XX 18-FEB-2002 (first entry)

XX DE Novel human diagnostic protein #21961.

XX KW Human; chromosome mapping; gene mapping; gene therapy; forensic;

KW food supplement; medical imaging; diagnostic; genetic disorder.

```
XX Homo sapiens.
OS WO200175067-A2.
PN 11-OCT-2001.
PD 30-MAR-2001; 2001WO-US08631.
PF 31-MAR-2000; 2000US-0540217.
PR 23-AUG-2000; 2000US-0649167.
PX (HYSE-) HYSEQ INC.
PA Drmanac RT, Liu C, Tang YT;
XX WPI; 2001-639362/73.
XX N-PSDB; AAS86157.
XX New isolated polynucleotide and encoded polypeptides, useful in
diagnostics, forensics, gene mapping, identification of mutations
responsible for genetic disorders or other traits and to assess
biodiversity
XX Claim 20; SEQ ID NO 52329; 103pp; English.
XX The invention relates to isolated polynucleotide (I) and
polypeptide (II) sequences. (I) is useful as hybridisation probes,
polymerase chain reaction (PCR) primers, oligomers, and for chromosome
and gene mapping, and in recombinant production of (II). The
polynucleotides are also used in diagnostics as expressed sequence tags
for identifying expressed genes. (I) is useful in gene therapy techniques
to restore normal activity of (II) or to treat disease states involving
(II). (II) is useful for generating antibodies against it, detecting or
quantitating a polypeptide in tissue, as molecular weight markers and as
a food supplement. (II) and its binding partners are useful in medical
imaging of sites expressing (II). (I) and (II) are useful for treating
disorders involving aberrant protein expression or biological activity.
The polypeptide and polynucleotide sequences have applications in
diagnostics, forensics, gene mapping, identification of mutations
responsible for genetic disorders or other traits to assess biodiversity
and to produce other types of data and products dependent on DNA and
amino acid sequences. ABG00010-ABG30377 represent novel human
diagnostic amino acid sequences of the invention.
XX Note: The sequence data for this patent did not appear in the printed
specification, but was obtained in electronic format directly from WIPO
at ftp.wipo.int/pub/published_pct_sequences.
XX SQ Sequence 575 AA;
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# alignment\_scores:

Quality: 105.00 Length: 519  
Ratio: 0.515 Gaps: 24  
Percent Similarity: 39.306 Percent Identity: 19.653

# alignment\_block:

US-09-303-518D-131 x ABG21970 ..

Align seq 1/1 to: ABG21970 from: 1 to: 575

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31 CCCATCGCGGGCAGACCGGAGCAAGTCATTATGACGGCCCGCCATTAC 80
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46 ProGlySerGlyLysSerProProValAlaSerGlyGlyProAla..... 60
81 CGAAGTCGCGTTCGTTGGCGAAGAAATATGTCGCATCGCGCCCTCGATGA 130
   :|||
61 .....ArgAlaLysProHisArgA 67
131 AAATCAAGGAAGGTGAAGCGCTCAAAAAGCCCAAGTGTCTTTGAAGAC 180
   :||| :|||
67 laValHisValSerProAlaProSerGlyGly..... 77
```









PI Daggett LP, Lu C;  
XX WPI; 2000-578607/54.  
DR N-PSDB; AAA95032.  
XX  
XX Novel DNA fragment encoding human N-methyl-D-aspartate receptor subunit  
PT for identifying mutations and for developing drugs against various  
PT disease states  
XX  
XX Example 3; column 255-264; 205pp; English.  
XX  
XX The present sequence is a subunit (designated NMDAR1A) of the human  
CC N-methyl-D-aspartate (NMDA) receptor. This is an ionotropic glutamate  
CC receptor which contains cation-specific ligand-gated ion channels. The  
CC protein and its coding sequence can be used in disease diagnosis and in  
CC research to identify other, similar proteins. They can also be used as  
CC probes, for example in genetic screening, and in drug screening, as well  
CC as enabling the production of animal disease models.  
XX  
XX Sequence 1232 AA;

## alignment\_scores:

Quality: 105.00 Length: 483  
Ratio: 0.593 Gaps: 28  
Percent Similarity: 36.646 Percent Identity: 23.602

## alignment\_block:

US-09-303-518D-131 x AAB26239 ..

Align seg 1/1 to: AAB26239 from: 1 to: 1232

26 ATCTGCCATCGCGGAGCGGAGCAAGTCATTATGACGGCCGCGCC 75  
856 LeuAlaProSerArgIleGlyValAlaAlaValArgProHisArgPR 872  
76 ATTACGAAGTCGGCTGCTGGCGAAGAATATGTCGCATCGCCCGCTC 125  
872 oProAlaArgProArgGlyLeuAlaPro...AlaHisAlaCysProPro. 887  
126 GATGAATCAAGGAGGTGAAGCGGTCAAAAAGG...CCAAGTGTCT 172  
888 .....ProThrArgPro.GlnSerArgAlaProArgAlaGl'899  
173 TTGAAGACAAAGAATCGCGGTAGTATTACTGCGCGCTTCAGGC 222  
899 yAspArgGlnThrGlyValAlaArgLeu...CysAlaGlyLeuArg. 914  
223 AAAATCGCGCTATTACCGTGGCGGAAAGCGCGT...ACTTCAGTCAGT 269  
915 ..SerProArgAlaAlaProArgArgGlyArgProCysProThrSer 930  
270 CGTGATTCGGTGAAGCAACGACGCAATCGAGTTCGAACGCTACGTAC 319  
931 ProGluCysArgAlaAlaGlnProGlyArgArgGlyArgCysGlyPR 947  
320 CTGAAGCGCTGCAAAATAGCAGCAAGAAAGTGCAGCGCAACCTGATT 369  
947 oGlyThrAlaGlyThrSerArgProProSerGlyPro..... 960  
370 CAATCAGGCTTATGACTCGCTTCGCACCCCTCCGTTCAACCAATCCC 419  
960 ..... 960  
420 TGGCGTAGATCGGAGCGGCTTCGCCATCTCTCAATGCGATGGACACCA 469  
960 ..... 960  
470 ATCCGCTGGCTCGCGACCCCTACCGTCATCATCAAGAAGCGCGCAAGAC 519  
961 .....CysArgPro.....ArgAlaV 966  
520 TTCAACGCGCGCTGTGTGATTGAGCCGCTGACCGAAGCTAAATCCA 569

966 alThrAlaProPhe...LeuGluProThrAspProAla...AlaPro 980  
570 TGTGTGTAAGCAGCAGCGGAGCGGCTGCTGAAATGC...TGCCA 616  
981 SerSerArgSerSerArgSerProArgSerTrpArgThrCysArgCysSe 997  
617 ATATCGAAACACATGAATTTGG.....CGGCCGCGATCC..... 650  
997 rValArgSerSer.....TrpProGlyGlyArgProCys\*\*\*ThrArgP 1012  
651 .....TGCGGCTTGTAGTGG 665  
1012 roGlyProGlyAlaArgAlaArgValThrLeuProCysProAlaProTrp 1028  
666 CAGGCACATTCATTCATCGACCGAGTCGCGCGAATAAAACCGTGTGA 715  
1028 ..... 1028  
716 CCATCAATTATCAAGACGTGATTGCTATCGGACGTTTGTCTGAACAGGC 765  
1029 .....ProArgP 1031  
766 CGTCTGAATACGAGCGCGCTGTTGCTTGGCGGCTCCAAGTCAACAA 815  
1031 roSerLeuGlyProAlaArgCysProLeuGlyAlaProAlaProAla 1047  
816 ACCGCGCTTGTGCGTACCGTGTGGTGGTGGAGGTTCTCAACTACCG 865  
1048 ProAlaProThrAlaThrArg.....ProAlaGlyAlaTrpAr 1060  
866 CCGCGCAATTTGTTGACGCGGACAAACCGCGTATTTCGGTTCGGTATTG 915  
1060 gAlGArg.....S 1063  
916 AAGCTTGGGATTCACAAAGCGCGCATGATTATTTGGGACGCTACCACAA 965  
1063 erArgCysAlaCys..... 1067  
966 TCAGATTTCGTTATCGAAGAGCGCGCAGCAAGAGCTGTTTCGGCTGGG 1015  
1068 .....ArgSerThrGlyArgProAlaArgAlaSerArgGlnG 1081  
1016 T..... 1016  
1081 yProProGlySerThrAspSerThrSerAlaCysThrProThrPro 1098  
1017 ..TGCGCGCAGCGCGCAAAATCTCC.....ATCAC 1046  
1098 hrCysHisCysAlaGlyGlyLeu.SerValLeuThrPheHisProValTh 1114  
1047 GCGCACCACTCTCGGCCATTTCTCTAAACAAACAACTCTCAAGTTCACGA 1096  
1114 rAlaThrAlaProGlySerProAlaProGlyGlyLeuTrpGlyThrAlaA 1131  
1097 CAGCGCTCAACGCGCGCGCATGTCGATCGGCACCTTATGAG 1146  
1131 laGlyLeuTrpGlyTrpAlaGlnAlaThrGluThrValGlyAspTrpThr 1147  
1147 CGC.....GTAATGCGTTGGACATCTCCCTACCTTGTCTTTGCGCGA 1190  
1148 ArgSerAlaVal\*\*ProValGlyArgLysAlaSer.....ArgAs 1161  
1191 TTTAATCGTCGGGATACCGACGCGGAGGCTTTGGTGTCTTGGTGAAT 1240  
1161 pProAlaProGlyAsp.....Gly.SerProVal 1170  
1241 TGGCAGGAGAGACCTCGCTTGTGACAGCTTCGTCTGCCCGGCAATAC 1290  
1171 TrpSerGlnLysCysGluLeuSerAlaThrGlnAla.ProSerGlnLeuA 1187  
1291 GAATACGCGCGCT.....GTTGCGCAAAAGTGTGGA 1322  
||||| ||| :|||

Mon Jul 1 09:25:33 2002

us-09-303-518d-131.rag

Page 57

1187 spSerLeuProAlaThrValArgVallysArgGlnAlaGly 1200

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